

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:20:48 ; Search time 307.385 Seconds

(without alignments)
10509.347 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111

Sequence: 1 atggttgatctcttgcctt.....gcctgagtgctgcttact 111

Scoring table: OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 2054640 seqs, 14551402878 residues

Word size : 0

Minimum number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

```

GenEmbl:*
1: gb_ba:*
2: gb_htg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vl:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pl:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vl:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htg_mus:*
34: em_htg_pin:*
35: em_htg_rod:*
36: em_htg_mam:*
37: em_htg_vrt:*
38: em_sy:*
39: em_htgo_hum:*
40: em_htgo_mus:*
41: em_htgo_other:*

```

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	111	100.0	169620	2	AC012674	AC012674 Homo sapi
2	48	43.2	143372	9	AL137847	AL137847 Human DNA
3	40	36.0	123779	30	AC021025	Ac021025 Homo sapi
4	40	36.0	128118	2	AC076969	AC076969 Homo sapi
5	40	36.0	128583	9	AC121249	AC121249 Homo sapi
6	40	36.0	148290	9	AC107979	AC107979 Homo sapi
7	40	36.0	165649	9	AC103996	AC103996 Homo sapi
8	40	36.0	178650	9	AC104303	AC104303 Homo sapi
9	40	36.0	192826	9	AC090762	AC090762 Homo sapi
10	39	35.1	32918	2	AC007445	AC007445 Homo sapi
11	39	35.1	38936	9	AL135817	AL135817 Human DNA
12	39	35.1	124271	2	AC025179	AC025179 Homo sapi
13	39	35.1	146671	9	AC008814	AC008814 Homo sapi
14	39	35.1	159747	2	AP001019	AP001019 Homo sapi
15	39	35.1	162740	2	AC034249	AC034249 Homo sapi
16	39	35.1	169772	9	AC069538	AC069538 Homo sapi
17	39	35.1	175466	9	AL607077	AL607077 Human DNA
18	38	34.2	110000	2	AL691517_2	Continuation (3 of
19	38	34.2	166706	9	AC068618	AC068618 Homo sapi
20	38	34.2	207408	2	AC087283	AC087283 Homo sapi
21	38	34.2	207548	9	AL591491	AL591491 Human DNA
22	36	32.4	121720	9	AL591491	AL591491 Human DNA
23	35	31.5	153940	9	AC022294	AC022294 Homo sapi
24	35	31.5	325069	2	AC079737	AC079737 Homo sapi
25	34	30.6	99577	9	AC026324	AC026324 Homo sapi
26	34	30.6	108040	2	AC068150	AC068150 Homo sapi
27	34	30.6	134760	9	AC099484	AC099484 Homo sapi
28	34	30.6	146059	9	AC019030	AC019030 Homo sapi
29	34	30.6	172206	9	AC092119	AC092119 Homo sapi
30	34	30.6	172567	2	AC015493	AC015493 Homo sapi
31	34	30.6	173166	9	AC092375	AC092375 Homo sapi
32	34	30.6	273807	2	AC025421	AC025421 Homo sapi
33	34	30.6	316296	2	AC092285	AC092285 Homo sapi
34	33	29.7	33458	9	HS0160101	AL109656 Human DNA
35	33	29.7	49616	9	AL365267	AL365267 Human DNA
36	33	29.7	77743	9	HSTCRBV	U03115 Human V bet
37	33	29.7	153788	2	AC023971	AC023971 Homo sapi
38	33	29.7	166434	9	AC020717	AC020717 Homo sapi
39	33	29.7	215422	9	U66060	U66060 Human germ
40	31	27.9	70313	2	AC016216	AC016216 Homo sapi
41	31	27.9	82225	9	HS253014	Z80771 Human DNA
42	31	27.9	164550	2	CNS01RHY	AL162633 Human chr
43	31	27.9	173706	9	AC022694	AC022694 Homo sapi
44	31	27.9	176418	9	AC090811	AC090811 Homo sapi
45	31	27.9	184759	9	AC022695	AC022695 Homo sapi

ALIGNMENTS

```

RESULT 1
LOCUS AC012674/c 169620 bp DNA linear HTG 07-SEP-2000
DEFINITION Homo sapiens chromosome 3 clone RPI-458H3, WORKING DRAFT SEQUENCE,
ACCESSION AC012674
VERSION AC012674.10 GI:9719580
KEYWORDS HTG: HTGS_PHASE1, HTGS_DRAFT.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 169620)
Muzny,D.M., Adams,C., Bailey,M., Barbara,J., Blankenburg,K.,
Bodola,B., Bouck,J., Bowie,S., Brooks,A., Buhay,C., Bunac,C.,

```

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

COMMENT

Burkett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C., David, R., Delgado, O., Deshazo, D., Ding, Y., Domah-Rashid, N., Dugan-Rocha, S., Durbin, K.J., Fernandez, C., Ferraguto, D., Forcum-Tansey, J., Frantz, P., Ganesh, R., Gorrell, J.H., Gorrell, L.L., Goullara, W., Harris, K., Hernandez, J., Hodgson, A., Hughes, M., Huelwa, C., Hosak, H., Jackson, L.E., Jackson, L., Jia, Y., Jones, M., Kelly, S., Kondejowski, N., Kong, Y., Kovar, C., Leal, B., Li, Z., Licharge, O., Liu, J., Liu, W., Logan, O., Lozano, R.J., Lu, J., Lucier, R., Martin, R., Martinez, C., McLeod, M.P., Mei, G., Morgan, M., Morris, S., Nash, S., Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S., Oswald, G., Parish, B., Paxton, A., Payton, B., Perez, L., Pu, L.L., Quinn, M., Reiter, D., Rives, M., Samuel, S., Say, J., Scherer, S., Shah, E., Shen, H., Simon, M., Sparks, A., Stamps, A., Sugeng, R., Taber, P., Taylor, T., Vasquez, L., Vinson, R., Vo, O., Webb, M., Wellington, S., Weinstein, G., Weinstein, I.R., Williamson, A., Worley, K., Wren, J., Wrensford, G., Yu, W., Zhou, X., Nelson, D., and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 169620)
Worley, K.C.

Direct Submission
Submitted (03-NOV-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Aug 7, 2000 this sequence version replaced gi:8705345.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HMO3
Center clone name: RPI-4558H3

----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 139025 bases at least Q40
Consensus quality: 154842 bases at least Q30
Consensus quality: 159725 bases at least Q20
Estimated insert size: 162720; sum-of-contigs estimation
Estimated insert size: 171608; agarose-fp estimation
Quality coverage: 3.9x in Q20 bases; agarose-fp estimation
Quality coverage: 4.1x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
(see <http://www.hgsc.bcm.tmc.edu/docs/genbank/draft.data.html>).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1
28689: contig of 28689 bp in length
28789: gap of unknown length
28790: contig of 22043 bp in length
50833: gap of unknown length
50932: gap of unknown length
50933: contig of 18212 bp in length
69144: gap of unknown length
69244: contig of 14960 bp in length
84205: gap of unknown length
84304: contig of 10363 bp in length
84305: gap of unknown length
94667: gap of unknown length
94767: contig of 12494 bp in length
10768: gap of unknown length
10769: contig of 10189 bp in length
107362: gap of unknown length
117550: contig of 10189 bp in length
117551: gap of unknown length
117650: contig of 9289 bp in length
126939: gap of unknown length
127038: contig of 8001 bp in length
127040: gap of unknown length
135040: contig of 8001 bp in length
135140: gap of unknown length
141639: contig of 6499 bp in length
135141

FEATURES
source
1. 169620
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RPI-4558H3"

BASE COUNT 52024 a 33180 c 32128 g 50322 t 1966 others

Query Match 100.0%; Score 111; DB 2; Length 169620;
Best Local Similarity 100.0%; Pred No. 2.3e-53;
Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 1 ATGGGTGATCTTTTGGCTTCAGGATCTTTTCATCTTTCAGGAGACTTTCGGGCCG 60
Db 87441 ATGGGTGATCTTTTGGCTTCAGGATCTTTTCATCTTTCAGGAGACTTTCGGGCCG 87382

OY 61 GAGATGTAACCTCCGGCTCTGTGTGCTGAGTGGCTGCTACT 111
Db 87381 GAGATGTAACCTCCGGCTCTGTGTGCTGAGTGGCTGCTACT 87331

RESULT 2
AL137847/c
LOCUS
DEFINITION Human DNA sequence from clone RPI1-439K3 on chromosome 9g422.2-31.1,
complete sequence.
ACCESSION AL137847 143372 bp DNA linear PRI 16-NOV-2001
VERSION AL137847
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
JOURNAL 1 (bases 1 to 143372)
Direct Submission

COMMENT
Submitted (16-NOV-2001) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk
On Nov 17, 2001 this sequence version replaced gi:16408610.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.

This sequence was either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >= 30): an attempt was made to resolve all sequencing problems, such
as compressions and repeats. All regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em, EMBL; SW,
SWISSPROT; Tr, TrEMBL; Wp, WormPeP; Information on the WormPeP

Matches 40: Conservative 0: Mismatches 0: Indels 0: Gaps 0:

QY 69 AAAACCTCGGCTCTGTGTGCTGCTAGTGGCTCTCT 108
DB 3083 AAAACCTCGGCTCTGTGTGCTGCTAGTGGCTCTCT 3044

RESULT 4

AC076969

LOCUS

DEFINITION

AC076969 128118 bp DNA linear HMG 15-OCT-2001
Homo sapiens chromosome 3 clone RP11-79K12, WORKING DRAFT SEQUENCE,
14 unordered pieces.

ACCESSION

AC076969

GI:16117967

VERSION

HTG: HTGS_PHASE1, HTGS_DRAFT.

KEYWORDS

Homo sapiens.

SOURCE

Homo sapiens

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE

1 (bases 1 to 128118)

AUTHORS

Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,

Alstrooms,S.L., Amaralunge,H.C., Are,J.R., Banks,T., Barabara,J.,

Benton,J., Bileva,M., Brown,E., Brown,M., Bryant,N.P., Bunay,C.,

Bowling,S., Burrell,C., Burrell,K.L., Byrd,N.C., Caron,T.F.,

Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,R.,

Chen,Z., Chowdhry,I., Christopoulos,C., Cleveland,C.D., Cox,C.,

Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C.,

Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O.,

Dem,A.L., Ding,Y., Dinh,H.H., Doultwaite,K.J., Drepper,H.,

Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C.,

Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J.,

Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T.,

Gara-N., Giller,R., Gorrell,J.H., Guevara,M., Gunaratne,P., Hale,S.,

Hamilton,K., Harris,C., Harris,K., Hart,M., Haylak,P., Hawes,A.,

Hernandez,J., Hernandez,O., Hodgson,A., Hogues,M., Holloway,C.,

Hollins,B., Homsi,F., Howard,S., Hubert,J., Huijy,S., Hume,J.,

Jaccobson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolliffe,S.,

Joudan,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvath,J.,

Kovar,C., Kralovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C.,

Lewis,L., Li,J., Li,Z., Licharge,O., Lieu,C., Liu,J., Liu,W.,

Lousaged,H., Lozano,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R.,

Ma,J., Maheshwari,M., Mapa,P., Martin,R., Martindale,A.,

Martinez,E., Massey,E., Mawhinney,E., McLeod,M.P., Meador,M.,

Mei,G., Metzger,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K.,

Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Newton,N.,

Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokweto,S.,

Ogun,M., Okunolu,G., Ogunye,N., Oviedo,R., Pace,A., Payton,B.,

Peery,J., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L.L.,

Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojebokan,I., Rolfe,M.,

Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shoostari,N.,

Slisson,I., Sodergren,E., Sonake,T., Sparks,A., Stanley,H.,

Stone,H., Sutton,A., Swatek,A., Taber,P., Tamerisa,A., Tamerisa,K.,

Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N.,

Thomas,S., Usmani,K., Vasquez,L., Vera,V., Villalob,D., Vinson,R.,

Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C.,

Watlington,S., Williams,G., Williamson,A., Wiczysk,R., Woodson,S.,

Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,

Weinstock,G. and Gibbs,R.

Unpublished

Direct Submission

2 (bases 1 to 128118)

Mortley,K.C.

Direct Submission

Submitted (01-AUG-2000) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

On Oct 14, 2001 this sequence version replaced gi:10047573.

Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu

Project Information

Center project name: HBRU

Center clone name: RP11-79K12

Sequencing statistics

Sequencing vector: M13: 108821

Assembly program: Phrap: version 0.990329

Consensus quality: 11307 bases at least 040

Consensus quality: 12194 bases at least 020

Estimated insert size: 122854; sum-of-coverage estimation

Quality coverage: 0x in 020 bases; agarose-gel estimation

Quality coverage: 4x in 020 bases; sum-of-coverage estimation

NOTE: Estimated insert size may differ from sequence length

(see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).

NOTE: This is a 'working draft' sequence. It currently

consists of 14 contigs. The true order of the pieces is

is not known and their order in this sequence record is

arbitrary. Gaps between the contigs are represented as

runs of N, but the exact sizes of the gaps are unknown.

This record will be updated with the finished sequence

as soon as it is available and the accession number will

be preserved.

1 20354: contig of 20354 bp in length

20355 20454: gap of unknown length

20455 33071: contig of 12617 bp in length

33072 33171: gap of unknown length

33172 46935: contig of 13764 bp in length

46936 47036: gap of unknown length

47036 57066: contig of 10031 bp in length

57067 57166: gap of unknown length

57167 70378: contig of 13212 bp in length

70379 70479: gap of unknown length

70479 81786: contig of 11308 bp in length

81787 81886: gap of unknown length

81887 91428: contig of 9542 bp in length

91429 91528: gap of unknown length

91528 100455: contig of 8927 bp in length

100456 100555: gap of unknown length

100556 107879: contig of 7324 bp in length

107880 107979: gap of unknown length

107980 114149: contig of 6170 bp in length

114150 114249: gap of unknown length

114250 120186: contig of 5937 bp in length

120187 120286: gap of unknown length

120287 123336: contig of 2950 bp in length

123337 123337: gap of unknown length

123337 125606: contig of 2270 bp in length

125607 125706: gap of unknown length

125707 128118: contig of 2412 bp in length.

Location/Qualifiers

1. 128118

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="3"

/clone="RP11-79K12"

BASE COUNT 35433 a 25053 c 23792 g 42367 t 1473 others

FEATURES

Source

Query Match

Best Local Similarity

Matches 40: Conservative

QY 69 AAAACCTCGGCTCTGTGTGCTGCTAGTGGCTCTCT 108

DB 55106 AAAACCTCGGCTCTGTGTGCTGCTAGTGGCTCTCT 55145

RESULT 5

AC121249/c

LOCUS

DEFINITION

AC121249 128583 bp DNA linear PRI 01-JUN-2002
Homo sapiens chromosome 3 clone RP11-79K17, complete sequence.

ACCESSION AC121249 AC021025
VERSION AC121249.1 GI:20806313
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
1 (bases 1 to 128583)
Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Saenphimmachak, C., Phelps, K.A., Buckley, D., Kibukawa, M., Raymond, C.
and Haugen, E.D.
Direct Submission
Unpublished
2 (bases 1 to 128583)
Kaul, R.K., Olson, M.V., Raymond, C. and Haugen, E.D.
Direct Submission
Submitted (16-MAY-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
3 (bases 1 to 128583)
Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Saenphimmachak, C., Phelps, K.A., Buckley, D., Kibukawa, M., Raymond, C.
and Haugen, E.D.
Direct Submission
Submitted (01-JUN-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
On May 16, 2002 this sequence version replaced gi:9719675.

Genome Center
Center: University of Washington Genome Center
Center Code: UMGc
Web site: http://www.genome.washington.edu
Contact: uwgchgs@u.washington.edu
Drafting Center: BCM

Project Information
Center project name: chr-3
Center clone name: RP11-79K17 (bc0196)

Summary Statistics: 45% of reads
Sequencing vector: plasmid; 55% of reads
Sequencing vector: M13; L08821; 5% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 128548 bases at least Q40
Consensus quality: 128580 bases at least Q30
Consensus quality: 128581 bases at least Q20
Insert size: 128583; sum-of-contigs
Quality coverage: 13.4x in Q20 bases; sum-of-contigs

Overlapping Sequences:
5' : RP11-147N17 (UMGC:bc0267) AC104300, 22116-bp overlap
3' : RP11-391P4 (UMGC:bc0402) AC104303, 114502-bp overlap

Sequence Quality Assessment:
This entry has been annotated with sequence quality
estimates computed by the Phrap assembly program.
All manually edited bases have been reduced to quality zero.
Quality levels above 40 are expected to have less than
1 error in 10,000 bp.
Base-by-base quality values are not generally visible from the
Genbank flat file format but are available as part
of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted:
all regions were either double-stranded or sequenced with an
alternate chemistry or covered by high quality data (i.e., Phred
quality >= 30); an attempt was made to resolve all sequencing
problems, such as compressions and repeats; all regions were
covered by at least one plasmid subclone or more than one M13
subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:
This sequence has been validated by Multiple Complete Digest
fingerprinting. Comparison of the experimentally derived digest
fragments with sequence-predicted fragments is given below.
The electronically-digested sequence consists of both insert and

vector, in order to accurately represent the entire circular BAC.
Small fragments below a variable cutoff (approximately 400-800 bp)
are not resolved in the fingerprint and hence do not appear
in the table. There are no significant remaining discrepancies
between the experimental and predicted values. Uniquely ordered
fragments are separated by dashed lines.

NsII		BgIII		EcoRI	
SeqDerMap	FingerPrint	SeqDerMap	FingerPrint	SeqDerMap	FingerPrint
12049	12136	4077	4013	8696	8902
579	<800	2067	2145	6	<800
1056	1065	9698	9932	1323	1337
13085	13126	3729	3734	1024	961
10944	10836	1632	1616	8421	8408
581	<800	8069	8139	1114	1126
882	887	1507	1485	3538	3534
8435	8467	11939	11871	6051	6022
3924	4103	133	<800	5129	5048
9842	9759	2135	2145	58	<800
4401	4377	78	<800	1895	1903
4458	4377	5873	5861	799	<800
4587	4522	896	902	888	885
6048	5947	3312	3356	931	885
1395	1385	1505	1485	859	885
3059	3071	6694	6716	3934	3936
3911	3861	10032	9932	1065	1058
138	<800	2168	2145	2629	2723
50	<800	1794	1785	859	885
582	<800	227	<800	6855	6888
2225	2229	4002	4013	7821	7819
984	967	2714	2726	7210	7201
887	887	5628	5624	6992	6888
1042	1065	743	758	1204	1224
2783	2808	3045	2963	2722	2723
7074	7158	1288	1262	321	<800
4087	3861	967	973	1066	1058
7976	7926	461	<800	470	<800
1233	1199	4649	4618	3055	3072
3875	3861	5289	5417	10961	10848
27	<800	1098	1095	6725	6888

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence.submissions@genome.wi.mit.edu

Project Information
Center project name: I24533
Center clone name: 3049_M_7

REFERENCE
AUTHORS
TITLE
JOURNAL
Submitted (01-DEC-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 165649)
Birtten,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhalter,B., Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A., Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Fero,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamt,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C., McCarthy,M., Meldrim,J., Menus,L., Mhova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Roy,A., Schauer,S., Schuback,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J., Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (26-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
4 (bases 1 to 165649)
Birtten,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhalter,B., Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A., Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Fero,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamt,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C., McCarthy,M., Meldrim,J., Menus,L., Mhova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Roy,A., Schauer,S., Schuback,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J., Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (01-JUL-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jul 1, 2002 this sequence version replaced gi:21592191.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
----- Project Information
Center project name: L21917
Center clone name: 76_E_17

FEATURES
source
Location/Qualifiers
1. 165649
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15"
/clone="RP11-76E17"
/clone_1kb="RPC1-11 Human Male BAC"
repeat_region
complement(2..865)
/rpt_family="L1PA13"
repeat_region
complement(864..894)
repeat_region
complement(895..1248)
/rpt_family="TRHE1A"
complement(1249..1504)
/rpt_family="L1PA13"
1505..1626
/rpt_family="Alusx"
/rpt_family="Alusx"
1627..1659
/rpt_family="(CAA)n"
/rpt_family="(CAA)n"
1660..1831
/rpt_family="Alusx"
complement(1832..3132)
/rpt_family="L1PA13"
complement(3140..3699)
/rpt_family="L1M4C"
complement(4461..4755)
/rpt_family="Alusx"
5027..5397
/rpt_family="L2"
complement(5650..6330)
/rpt_family="L1ME1"
complement(6343..6460)
/rpt_family="L1ME1"
7214..7319
/rpt_family="L1MC3"
7354..7655
/rpt_family="L1MC3"
7659..7743
/rpt_family="L1MC3"
complement(7745..7843)
/rpt_family="Alusp/q"
7844..14156
/rpt_family="L1PA10"
14198..14328
/rpt_family="Aluy"
14340..14503
/rpt_family="(TA)n"
14507..15032
/rpt_family="L1MC3"
15036..15220
/rpt_family="L1MC3"
complement(15218..16449)
/rpt_family="L1PA4"
16450..18154
/rpt_family="L1PA4"
18155..19210
/rpt_family="L1MC3"
19211..19266
/rpt_family="(TA)n"
19267..19372
/rpt_family="L1MC3"
19401..19476
/rpt_family="(TTATA)n"
19483..19546
/rpt_family="(CATATA)n"
19596..19625
/rpt_family="AT-rich"
19696..19750
/rpt_family="GA-rich"
19752..19920
/rpt_family="L1MD3"
19994..20102
/rpt_family="L2"
20049..20114
/note="single clone coverage"
20485..20655
/rpt_family="MIR3"
complement(21285..21441)
/rpt_family="MIR"
21496..21717
/rpt_family="L2"
complement(21720..21796)
/rpt_family="MIR"

repeat_region	complement(21943. .22131)
/rpt_family="MIR"	23082. .23195
repeat_region	/rpt_family="r2"
repeat_region	23198. .23348
/rpt_family="GA-rich"	23267. .23333
repeat_region	/rpt_family="(CAT)n"
repeat_region	complement(24465. .24833)
/rpt_family="MTA1a2"	26142. .26334
repeat_region	/rpt_family="MIR"
repeat_region	complement(27659. .27811)
/rpt_family="MIR"	28880
repeat_region	/rpt_family="MIR"
repeat_region	complement(29026. .29269)
/rpt_family="MIR"	complement(29673. .29706)
repeat_region	/rpt_family="MTB"

```

every Match      36.0%; Score 40; DB 9; Length 165649;
Best Local Similarity 100.0%; Pred. No. 4.3e-12;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

07 68 TAAACCTCTGGGTCTCTGTGTGCTAGTGGCTGCTC 107
 |||||
 Db 8110 TAAACCTCTGGGTCTCTGTGTGCTGAGTGGCTGCTC 807

RESULT	8
LOCUS	AC104303
DEFINITION	Homo sapiens chromosome 3 clone RP11-391P4, complete sequence.
ACCSSION	AC104303.AC064830
VERSION	AC104303.2 GI:18874945
KEYWORDS	HTG.
SOURCE	Homo sapiens.
ORGANISM	Homo sapiens

REFERENCE 1 (Phases 1 to 178650)
AUTHORS Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Seephimmachak, C., Phelps, K.A., Raymond, C. and Haugen, E.D.
TITLE Direct Submission
JOURNAL Unpublished
REFERENCE 2 (Phases 1 to 178650)
AUTHORS Kaul, R.K., Olson, M.V., Raymond, C. and Haugen, E.D.
TITLE Direct Submission
JOURNAL Submitted (07-DEC-2001) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
3 (Phases 1 to 178650)
AUTHORS Kaul, R.K., Olson, M.V., Zhou, Y., James, R.A., Rouse, G., Wu, Z.,
Seephimmachak, C., Phelps, K.A., Raymond, C. and Haugen, E.D.
TITLE Direct Submission
JOURNAL Submitted (25-FEB-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
COMMENT On Feb 25, 2002 this sequence version replaced gi:17402782.

```

Center: University of Washington Genome Center
Center Code: UWGC
Web site: http://www.genome.washington.edu
Contact: uwgchugst@u.washington.edu
Drafting Center: UWGSC
-----
Project Information
Center project name: chr-3
Center clone name: RP11-391P4 (bc0402)
-----
Summary Statistics
Sequencing vector: unknown; 55% of reads
Sequencing method: plasmid; 45% of reads
Chemistry: dye-terminator ET; 89% of reads
Chemistry: dye-terminator Big Dye; 11% of reads
Assembly program: Phrap; version 0.990319

```

Consensus quality: 178494 bases at least Q40
Consensus quality: 178631 bases at least Q30
Consensus quality: 178650 bases at least Q20
Insert size: 178648; sum-of-contigs
Quality coverage: 8.0x in Q20 bases; sum-of-contigs

overlapping Sequences:
5': RP11-475023 (UMGC:bc0439) AC023346
3': RP11-79K12 AC076969

Sequence Quality Assessment:

This entry has been annotated with sequence quality estimates computed by the Phrap assembly program. All manually edited bases have been reduced to quality zero. Quality levels above 40 are expected to have less than 1 error in 10,000 bp. Base-by-base quality values are not generally visible from the Genbank flat file format but are available as part of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., Phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest fingerprinting. Comparison of the experimentally derived digest fragments with sequence-predicted fragments is given below. The electronically-digested sequence consists of both insert and vector, in order to accurately represent the entire circular BAC. Small fragments below a variable cutoff (approximately 400-800 bps) are not resolved in the fingerprint and hence do not appear in the table. There are no significant remaining discrepancies between the experimental and predicted values. Uniquely ordered fragments are separated by dashed lines.

Bq111

FCORI

HindIII

SedDerMap	FngPrnt	SedDerMap	FngPrnt	SedDerMap	FngPrnt
7050	7417	8696	8715	7163	7098
2067	2138	6	<800	6582	6518
13472	13264	3077	3083	512	<800
7644	7859	514	<800	449	<800
111	<800	4438	4377	7988	7884
798	783	8682	8715	11779	11698
1098	1109	1896	1876	472	<800
5288	5395	6724	6948	783	787
4649	4507	10961	10773	926	926
461	<800	3055	3083	2431	2538
967	982	470	<800	1015	1029
1288	1268	1066	1037	758	<800
3045	2923	321	<800	1550	1541
743	783	2722	2683	361	<800

	5628	5682	1204	1300	763	<800
	2714	2699	6992	7281	2975	3002
	4002	3900	7210	7830	3299	3468
	227	<800	7821	8101	1907	1894
	1794	1784	6855	6948	4020	4220
	2168	2138	859	857	5158	5126
	10032	9744	2629	2683	457	<800
	6694	6810	1065	1037	4321	4220
	1505	1491	3934	3965	887	926
	3312	3304	859	857	180	<800
	896	912	931	935	4969	4887
	5873	5938	858	857	4755	4674
	78	<800	799	857	26	<800
	2135	2138	1895	1876	906	926
	133	<800	58	<800	2325	2346
	11939	11575	5129	5076	369	<800
	1507	1491	6051	6048	2523	2538
	8069	8181	3538	3537	1889	1894
	1632	1617	1114	1094	4437	4428
	3729	3678	8420	8391	373	<800
	5553	5395	1024	1037	24	<800
	358	<800	1333	1380	263	<800
	3764	3678	1502	1481	632	<800
	7022	6810	4538	4539	1258	1247
	8534	8821	1414	1380	1114	1083
	5206	5050	521	<800	1370	1320
	9052	9744	8346	8391	1262	1247
	1625	1617	39	<800	1546	1541
	749	783	2716	2683	459	<800
	10190	9744	2546	2683	135	<800
	3857	3795	1064	1037	3142	3183
	6144	6490	926	935	109	<800
	514	<800	5154	5076	7858	7884
	1364	1342	1975	1981	1891	1894
	669	<800	628	<800	723	<800
			2398	2399	6975	7098
			8148	8391	1878	1894

Query Match	36.0%;	Score 40;	DB 9;	Length 178650;
Best Local Similarity	100.0%;	Pred. No. 4.3e-12;		
Matches 40;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

QY 69 AAACATCCTGGGCTCTGTGTGTGCTGCTGAGTGGCTCTCT 108
|||||
Db 165449 AAACATCCTGGGCTCTGTGTGTGCTGCTGAGTGGCTCTCT 1654888

RESULT	9
LOCUS	AC090762/c
DEFINITION	Homo sapiens chromosome 15, clone RP11-387E8, complete sequence.
ACCESSION	AC090762
VERSION	AC090762.9
KEYWORDS	HTG.
SOURCE	Homo sapiens.
ORGANISM	Homo sapiens

REFERENCE	1 (bases 1 to 192826)
AUTHORS	Birren, B., Linton, L., Nussbaum, C. and Lander, E.
TITLE	Homo sapiens chromosome 15, clone RP11-387E8
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 192826)
AUTHORS	Birren, B., Linton, L., Nussbaum, C., Lander, E., Allen, N., Anderson, S.,

AUTHORS
Birtten, B., Litton, L., Nussbaum, C., Lander, E., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Boguslavsky, L., Bouhagalter, B., Brown, A.,
Cammarato, J., Campopiano, A., Choelely, Y., Colangelo, M., Collins, S.,
Collamore, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S.,
Dodge, S., Fato, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J.,
Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N.,
Hagos, B., Heatford, A., Horton, L., Hulme, W., Illey, I., Johnson, R.,
Jones, C., Karlzas, A., Larocque, K., Lamazates, R., Landers, T.,
Lehoczky, J., Levine, R., Liu, G., Maclean, C., MacDonald, P.,
Marquis, N., Matthews, C., McCarthey, M., McEwan, P., McKernan, K.,
McPheeters, R., Meldrum, J., Meneus, L., Mihova, T., Mlenka, V.,
Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C. H.,
O'Connor, I., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
Phunhkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R.,
Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Roselli, M.,
Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S., Severy, P.,
Sounguez, C., Spencer, B., Strange-Thomann, N., Stojanovic, N.,
Strauss, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J.,
Travers, M., Travis, N., Triggillo, J., Vassiliev, H., Viel, R., Vo, A.,
Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.

TITLE Direct Submission
JOURNAL Submitted (10-MAR-2001) Whitehead Institute/MIT Center for Genome
REFERENCE Research, 330 Charles Street, Cambridge, MA 02141, USA
AUTHORS 3 (bases 1 to 192826)
Birren, B., Linton, L., Nusbaum, C., Lander, E., All, A., Allen, N.,
Anderson, S., Barna, N., Bastien, Y., Boguslavsky, I., Boukhalter, B.,
Brown, A., Camarata, J., Campianno, A., Chang, J., Chazaro, B.,
Choehel, Y., Collangelo, M., Collins, S., Collymore, A., Cook, A.,
Cooke, P., Dearlano, K., Desai, K., Diaz, J. S., Dodge, S., Faro, S.,
Ferreira, P., Fitzhugh, W., Gage, D., Galagan, U., Galya, S.,
Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N.,
Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,
Kamat, A., Karats, A., Kells, C., Lacroque, K., Lamazeres, R.,

TITLE
JOURNAL
COMMENT

Landers, T., Lehoczy, J., Levine, R., Liu, G., MacLean, C.,
Maddonald, P., Major, J., Margus, N., Matthews, C., McCarthy, M.,
McKernan, P., McKernan, K., Meldrum, J., Meneus, L., Mihova, T.,
Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C.,
Notman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J.,
Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C.,
Retta, R., Riback, M., Riley, R., Rise, C., Rogov, P., Roman, J.,
Roselli, M., Roy, A., Santos, R., Schauer, S., Schuppach, R., Seaman, S.,
Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
Topham, K., Travers, M., Travis, N., Triggillo, J., Vassiliev, H.,
Vieth, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G.,
Zainoun, J., Zemdek, L., Zimmer, A. and Zody, M.

Submitted (28-FEB-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Feb 28, 2002 this sequence version replaced g1:18377189.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIRB
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence.submissions@genome.wi.mit.edu
Project Information
Center project name: L12392
Center clone name: 387_E_8

FEATURES

Source

Location/Qualifiers
1. 192826
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15"
/clone="RP11-387B8"
/clone_lib="RPCT-11 Human Male BAC"
repeat_region
complement(864..1222)
/rpt_family="L1PB3"
1223..1252
/rpt_family="(TAGC)n"
complement(1253..1492)
repeat_region
/rpt_family="L1PB3"
4287..4576
/rpt_family="AluJb"
4671..5034
/rpt_family="THE1B"
5130..5438
/rpt_family="AluSx"
5517..5662
/rpt_family="MIR"
5798..5827
/rpt_family="AT-rich"
5985..6114
/rpt_family="AluJb"
6227..6253
/rpt_family="(CA)n"
7329..7446
/rpt_family="MIR"
complement(8147..8452)
/rpt_family="AluJo"
8540..8639
/rpt_family="MER45"
complement(8832..8916)
/rpt_family="L2"
10391..10599
/rpt_family="MER3"
11688..11688
/rpt_family="L1MC/D"
12029..12078
/rpt_family="AT-rich"
12092..12447
/rpt_family="THE1C"

repeat_region complement(13616..13751)
/rpt_family="MIR3"
repeat_region complement(13958..14142)
/rpt_family="MIR"
repeat_region complement(15277..15553)
/rpt_family="MER8"
repeat_region complement(15811..16005)
/rpt_family="MIR"
repeat_region complement(16101..16440)
/rpt_family="L3"
16920..16958
/rpt_family="(TCCC)n"
repeat_region complement(17145..17444)
/rpt_family="AluSx"
repeat_region complement(18418..19953)
/rpt_family="L1MEC"
repeat_region complement(19978..20262)
/rpt_family="L1MEC"
repeat_region complement(20288..20794)
/rpt_family="L1MEC"
repeat_region complement(20822..21097)
/rpt_family="L1MEC"
repeat_region complement(21345..21743)
/rpt_family="L1MEC"
22599..23518
/rpt_family="L1MEC"
23527..23901
/rpt_family="L1MEC"
repeat_region complement(23927..24026)
/rpt_family="MSTR1"
repeat_region complement(24027..24256)
/rpt_family="MER30"
repeat_region complement(24257..24557)
/rpt_family="MSTR1"
24563..24594
/rpt_family="AT-rich"
repeat_region complement(25258..25639)
/rpt_family="L1MCc"
repeat_region complement(25688..25838)
/rpt_family="L1MCc"
26167..26506
/rpt_family="THE1B"
26746..27094
/rpt_family="Tiger2a"
27095..27184
/rpt_family="MADE1"
27185..27279
/rpt_family="Tiger2a"
28321..28363
/rpt_family="(TG)n"
29172..29333
/rpt_family="MIR"
30664..30898
/rpt_family="L1MB8"
31577..31598
/rpt_family="AT-rich"
32378..32475
/rpt_family="CT-rich"
34655..34960
/rpt_family="AluY"
repeat_region complement(35296..35389)
/rpt_family="MER5B"
repeat_region complement(35419..35517)
/rpt_family="L1MC4"
repeat_region complement(35518..35823)
/rpt_family="AluSx"
repeat_region complement(35824..36030)
/rpt_family="L1MC4"
36682..36758
/rpt_family="MER5A"
36975..37160
/rpt_family="MER5A"
repeat_region 38140..38353

```

repeat_region      /rpt_family="MER96B"
                   38633..38656
repeat_region      /rpt_family="(TTTA)n"
                   complement(38657..40056)
repeat_region      /rpt_family="LIPAS"
                   complement(40060..40250)

Query Match
Best Local Similarity 100.0%; Score 40; DB 9; Length 192826;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 68 TAAACTCTGGCTCTGTGTGTGCTGAGTGCTGCTC 107
      |||
Db 190946 TAAACTCTGGCTCTGTGTGCTGAGTGCTGCTC 190907

RESULT 10
AC007445      32918 bp      DNA      linear      HTG 30-JUN-2000
US            Homo sapiens chromosome 18 clone RP11-344B7 map 18, *** SEQUENCING
AC007445      IN PROGRESS *** 1 ordered piece.
AC007445      HTG: HTGS_PPHASE2.
KEYWORDS      Homo sapiens.
SOURCE        Homo sapiens
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE      1 (bases 1 to 32918)
AUTHORS        Birren,B., Linton,L., Nusbaum,C. and Lander,E.
JOURNAL        Homo sapiens chromosome 18, clone RP11-344B7
TITLE          Unpublished
COMMENT        2 (bases 1 to 32918)
              Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,M.,
              Baker,J., Baldwin,J., Barna,N., Beckey,R., Benn,J., Brown,A.,
              Castle,A., Cerny,J., Colangelo,M., Collins,S., Collymore,A.,
              Cooke,P., Dekrelano,K., Depayre,E., Devon,K., Dewar,K.,
              Donelan,L., Doyle,M., Ferrelta,P., FitzHugh,W., Forrest,C.,
              Funke,R., Gage,D., Galagan,J., Gardyna,S., Gilbert,D., Grant,G.,
              Hagos,B., Heaford,A., Horton,L., Howland,J.C., Jones,C., Kann,L.,
              Karats,A., Lehoczy,J., Lieu,C., Locke,K., Macdonald,P.,
              Marquis,N., McEwan,P., McGurk,A., McKernan,K., McLaughlin,J.,
              Meidrum,J., Molla,M., Morris,W., Morrow,J., Mychaleckyj,J.,
              Naylor,J., Niloff,M., O'Connor,T., O'Donnell,P., Pavlin,B.,
              Peterson,K., Pollara,V., Riley,R., Roberts,D., Roy,A., Severy,P.,
              Stange-Thomann,N., Stojanovic,N., Stone,C., Subramanian,A.,
              Testaye,S., Torruella-Miller,I., Vassiliev,H., Vo,A., Wagner,A.,
              Wheeler,J., Wu,X., Wyman,D., Ye,W.J. and Zody,M.
              Direct Submission
              Submitted (30-APR-1999) Whitehead Institute/MIT Center for Genome
              Research, 320 Charles Street, Cambridge, MA 02141, USA
              On Jun 30, 2000 this sequence version replaced g1:8705092.
              All repeats were identified using RepeatMasker:
              Smit, A.F.A. & Green, P. (1996-1997)
              http://ftp.genome.washington.edu/RM/RepeatMasker.html

              Genome Center
              Center: Whitehead Institute/ MIT Center for Genome Research
              Center code: WIBR
              Web site: http://www-seq.wi.mit.edu
              Contact: sequence_submissions@genome.wi.mit.edu
              ----- Project Information
              Center project name: L571
              Center clone name: 344_B-7

```

```

* NOTE: This is a 'working draft' sequence. It currently
* consists of 1 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submittor.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.

```

```

FEATURES
source
1 32918: contig of 32918 bp in length.
Location/Qualifiers
1..32918
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
/map="18"
/clone="RP11-344B7"
/clone_lib="RPC1-11 Human Male BAC"
BASE COUNT 9462 a 6493 c 6865 g 9853 t 245 others
ORIGIN

Query Match
Best Local Similarity 100.0%; Score 39; DB 2; Length 32918;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 60 GGAGTATGTAACCTCGTCTGTGTGTGCTGAG 98
      |||
Db 3535 GGAGTATGTAACCTCGTCTGTGTGTGCTGAG 3573

RESULT 11
AL358817      38936 bp      DNA      linear      PRI 06-OCT-2001
LOCUS         Human DNA sequence from clone RP11-399N22 on chromosome 10,
DEFINITION    complete sequence.
ACCESSION     AL358817
VERSION       AL358817.18 GI:15990637
KEYWORDS      HTG.
SOURCE        human.
ORGANISM      Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE      1 (bases 1 to 38936)
AUTHORS        Lovell,J.
JOURNAL        Direct Submission
              Submitted (06-OCT-2001) Sanger Centre, Hinxton, Cambridgeshire,
              CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
              On Oct 9, 2001 this sequence version replaced g1:14669268.
              requests: clonerequests@sanger.ac.uk
              During sequence assembly data is compared from overlapping clones.
              Where differences are found these are annotated as variations
              together with a note of the overlapping clone name. Note that the
              variation annotation may not be found in the sequence submission
              corresponding to the overlapping clone, as we submit sequences with
              only a small overlap as described above.
              This sequence was finished as follows unless otherwise noted: all
              regions were either double-stranded or sequenced with an alternate
              chemistry or covered by high quality data (i.e., phred quality >=
              30); an attempt was made to resolve all sequencing problems, such
              as compressions and repeats; all regions were covered by at least
              one plasmid subclone or more than one M13 subclone; and the
              assembly was confirmed by restriction digest. The following
              abbreviations are used to associate primary accession numbers given
              in the feature table with their source databases: Em:, EMBL; SW:,
              SWISSPROT; Tr:, TrEMBL; Wp:, WormPEP; Information on the WormPEP
              database can be found at
              http://www.sanger.ac.uk/Projects/C_elegans/wormpep
              This sequence
              was generated from part of bacterial clone contigs of human
              chromosome 10, constructed by the Sanger Centre Chromosome 10
              Mapping Group. Further information can be found at
              http://www.sanger.ac.uk/HGP/Chr10
              RP11-399N22 is from the library RPC1-11.2 constructed by the group
              of Pieter de Jong. For further details see
              http://www.chori.org/bacpac/home.htm
              VECTOR: pBAC3.6
              IMPORTANT: This sequence is not the entire insert of clone
              RP11-399N22. It may be shorter because we sequence overlapping
              sections only once, except for a short overlap.
              The true left end of clone RP11-43299 is at 36937 in this sequence.
              The true right end of clone RP11-91A1 is at 2000 in this sequence.
              Location/Qualifiers
              1..38936

```

```

FEATURES
source

```


/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="10"
/clone="RP11-399N22"
/clone_lib="RPCT-11.2"
/clone_lib="RPCT-11.2"
BASE COUNT 9315 a 9079 c 9111 g 11431 t
ORIGIN

Query Match 35.1%; Score 39; DB 9; Length 38936;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 60 GGAGTATGTAACCTCTGGTCTCTGTGTGCTGAG 98
|||||
Db 28896 GGAGTATGTAACCTCTGGTCTCTGTGTGCTGAG 28934

RESULT 12
AC025179 124271 bp DNA linear HTG 20-APR-2001
LOCUS Homo sapiens chromosome 5 clone CTD-2174B5, WORKING DRAFT SEQUENCE,
DEFINITION 8 unordered pieces.
AC025179
AC025179.4 GI:13699647
VERSION HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEFN.
KEYWORDS Homo sapiens.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 124271)
TITLE DOE Joint Genome Institute.
AUTHORS Sequencing of Human Chromosome 5
JOURNAL Unpublished
2 (bases 1 to 124271)
TITLE DOE Joint Genome Institute.
AUTHORS Direct Submission
JOURNAL Submitted (07-MAR-2000) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
On Apr 20, 2001 this sequence version replaced gi:7711794.
-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov

Project Information
Center Project Name: 694394
Center clone name: CITB-HL_2174B5

Summary Statistics
Consensus quality: 116609 bases at least Q40
Consensus quality: 120226 bases at least Q30
Consensus quality: 121274 bases at least Q20
Estimated insert size: 117160; agarose-fp estimation
Estimated insert size: 123571; sum-of-contigs estimation
Quality coverage: 6.46 in Q20 bases; agarose-fp estimation
Quality coverage: 6.12 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 8 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 1052: contig of 1052 bp in length
* 1053 1152: gap of unknown length
* 1153 3331: contig of 2179 bp in length
* 3332 3431: gap of unknown length
* 3432 7370: contig of 3939 bp in length
* 7371 7470: gap of unknown length
* 7471 15483: contig of 8013 bp in length
* 15484 15583: gap of unknown length
* 15584 24916: contig of 9333 bp in length

* 24917 25016: gap of unknown length
* 25017 39922: contig of 1490 bp in length
* 39923 40022: gap of unknown length
* 40023 68684: contig of 28662 bp in length
* 68685 68785: gap of unknown length
* 68785 124271: contig of 55487 bp in length.
Location/Qualifiers
1. 124271
source
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTD-2174B5"
/clone_lib="Caltech human BAC library D"
BASE COUNT 36863 a 24233 c 23599 g 38876 t 700 others
ORIGIN

Query Match 35.1%; Score 39; DB 2; Length 124271;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCTGGTCTCTGTGTGCTGAGT 99
|||||
Db 79646 GAGTATGTAACCTCTGGTCTCTGTGTGCTGAGT 79684

RESULT 13
AC008814 146671 bp DNA linear PRI 31-OCT-2001
LOCUS Homo sapiens chromosome 5 clone CTD-21171L12, complete sequence.
DEFINITION AC008814
AC008814
AC008814.6 GI:16554342
VERSION HTG.
KEYWORDS Homo sapiens.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 146671)
TITLE DOE Joint Genome Institute and Stanford Human Genome Center.
AUTHORS Direct Submission
JOURNAL Unpublished
2 (bases 1 to 146671)
TITLE DOE Joint Genome Institute.
AUTHORS Direct Submission
JOURNAL Submitted (03-AUG-1999) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
3 (bases 1 to 146671)
TITLE DOE Joint Genome Institute and Stanford Human Genome Center.
JOURNAL Direct Submission
Submitted (31-OCT-2001) DOE Joint Genome Institute, 2800 Mitchell
Drive, Walnut Creek, CA 94598, USA
On Oct 31, 2001 this sequence version replaced gi:15290309.
Draft Sequence Produced by DOE Joint Genome Institute
www.jgi.doe.gov
Finishing Completed at Stanford Human Genome Center
www.shgc.stanford.edu
Quality: Phrap Quality >=40 99.3% of Sequence;
Estimated Total Number of Errors is 0.7.
Location/Qualifiers
1. 146671
source
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTD-21171L12"
BASE COUNT 44951 a 28815 c 28434 g 44471 t
ORIGIN

Query Match 35.1%; Score 39; DB 9; Length 146671;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCTGGTCTCTGTGTGCTGAGT 99
|||||
Db 54480 GAGTATGTAACCTCTGGTCTCTGTGTGCTGAGT 54518

RESULT 14
AP001019/c
LOCUS Homo sapiens chromosome 18 clone RP11-752111 map 18p11.3, WORKING
DEFINITION DRAFT SEQUENCE, 24 unordered pieces.
ACCESSION AP001019
VERSION AP001019.2 GI:8117689
KEYWORDS HTG: HTGS, PHASE1: HTGS, DRAFT.
SOURCE Homo sapiens DNA, clone:RP11-752111.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 159747)
Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
Fujiyama, A., Yada, T., Totoki, Y., Matanabe, H. and Sakaki, Y.
Homo sapiens 159,747 genomic DNA of 18p11.3
2 (bases 1 to 159747)
Hattori, M., Ishii, K., Toyoda, A., Taylor, T.D., Hong-Seog, P.,
Fujiyama, A., Yada, T., Totoki, Y., Matanabe, H. and Sakaki, Y.
Direct Submission
Submitted (05-JAN-2000) Masahira Hattori, The Institute of Physical
and Chemical Research (RIKEN), Genomic Sciences Center (GSC);
Kitasato Univ., 1-15-1 Kitasato, Sagamihara, Kanagawa 228-8555,
Japan (E-mail:hattori@gscc.riken.go.jp,
URL:http://hgp.gscc.riken.go.jp/, Tel:81-42-778-9923,
Fax:81-42-778-9924)
On May 31, 2000 this sequence version replaced gi:6997769.
----- Genome Center
Center: RIKEN Genomic Sciences Center(GSC)
Center code: RIKEN
Web site: http://hgp.gscc.riken.go.jp/
Contact: hattori@gscc.riken.go.jp
----- Project Information
Center project name: Humdraft18
Center clone name: RP11-752111
----- Summary Statistics
Sequencing vector: PCR products; 100% of reads
Chemistry: Dye-terminator ET-amersham; 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 145356 bases at least Q40
Consensus quality: 152227 bases at least Q30
Consensus quality: 155843 bases at least Q20
Insert size: 157447; sum-of-contigs
Quality coverage: 4.51x in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of
24 contigs. The true order of the pieces is not known and their
order in this sequence record is arbitrary. Gaps between the
contigs are represented as runs N, but the exact sizes of the gaps
are unknown. This record will be updated with the finished sequence
as soon as it is available and the accession number will be
preserved
1 18444 contig of 18444 bp in length
18545 29631 contig of 11087 bp in length
29732 43143 contig of 13412 bp in length
43244 57170 contig of 13927 bp in length
57271 68240 contig of 10970 bp in length
68341 78134 contig of 9794 bp in length
78235 87873 contig of 9639 bp in length
87974 96197 contig of 8224 bp in length
96298 103976 contig of 7679 bp in length
104077 110726 contig of 6650 bp in length
110827 116866 contig of 5823 bp in length
116967 122789 contig of 5823 bp in length
122890 127666 contig of 4776 bp in length
127766 132618 contig of 4853 bp in length
132619 137718 contig of 500 bp in length
137719 136051 contig of 3333 bp in length
136052 136151 contig of 100 bp in length
136152 139749 contig of 3598 bp in length
139750 139849 contig of 100 bp in length
139850 142810 contig of 2861 bp in length
142811 142910 contig of 100 bp in length
142911 145855 contig of 2945 bp in length
145856 145955 contig of 100 bp in length
145956 148840 contig of 2885 bp in length
148841 148940 contig of 100 bp in length
148941 151243 contig of 2203 bp in length
151244 151343 contig of 100 bp in length
151344 153454 contig of 2111 bp in length
153455 153554 contig of 100 bp in length
153555 156134 contig of 2580 bp in length
156135 156234 contig of 100 bp in length
156235 158089 contig of 1555 bp in length
158090 158189 contig of 100 bp in length
158190 159747 contig of 1558 bp in length.

FEATURES
source
1..159747
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
/map="18p11.3"
/clone="RP11-752111"
1..18444
/note="assembly_fragment"
18545..29631
/note="assembly_fragment"

145956 148840 contig of 2885 bp in length
148941 151243 contig of 2303 bp in length
151344 153454 contig of 2111 bp in length
153555 156134 contig of 2580 bp in length
156235 158089 contig of 1855 bp in length
158190 159747 contig of 1558 bp in length
Sequence updated (26-May-2000)
* NOTE: This is a 'working draft' sequence. It currently
* consists of 24 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1 18444: contig of 18444 bp in length
18445 18544: gap of 100 bp
18545 29631: contig of 11087 bp in length
29632 29731: gap of 100 bp
29732 43143: contig of 13412 bp in length
43144 43243: gap of 100 bp
43244 57170: contig of 13927 bp in length
57171 57270: gap of 100 bp
57271 68240: contig of 10970 bp in length
68241 68340: gap of 100 bp
68341 78134: contig of 9794 bp in length
78135 78234: gap of 100 bp
78235 87873: contig of 9639 bp in length
87874 87973: gap of 100 bp
87974 96197: contig of 8224 bp in length
96198 96297: gap of 100 bp
96298 103976: contig of 7679 bp in length
103977 104076: gap of 100 bp
104077 110726: contig of 6650 bp in length
110727 110826: gap of 100 bp
110827 116866: contig of 6040 bp in length
116867 116966: gap of 100 bp
116967 122789: contig of 5823 bp in length
122790 122889: gap of 100 bp
122890 127665: contig of 4776 bp in length
127666 127765: gap of 100 bp
127766 132618: contig of 4853 bp in length
132619 137718: gap of 100 bp
137719 136051: contig of 3333 bp in length
136052 136151: gap of 100 bp
136152 139749: contig of 3598 bp in length
139750 139849: gap of 100 bp
139850 142810: contig of 2861 bp in length
142811 142910: gap of 100 bp
142911 145855: contig of 2945 bp in length
145856 145955: gap of 100 bp
145956 148840: contig of 2885 bp in length
148841 148940: gap of 100 bp
148941 151243: contig of 2203 bp in length
151244 151343: gap of 100 bp
151344 153454: contig of 2111 bp in length
153455 153554: gap of 100 bp
153555 156134: contig of 2580 bp in length
156135 156234: gap of 100 bp
156235 158089: contig of 1555 bp in length
158090 158189: gap of 100 bp
158190 159747: contig of 1558 bp in length.

FEATURES
source
1..159747
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
/map="18p11.3"
/clone="RP11-752111"
1..18444
/note="assembly_fragment"
18545..29631
/note="assembly_fragment"

```
misc-feature 29732..43143
              /note="assembly-fragment"
misc-feature 43244..57170
              /note="assembly-fragment clone_end:SP6 vector_side:left"
misc-feature 57271..68240
              /note="assembly-fragment"
misc-feature 68341..78134
              /note="assembly-fragment"
misc-feature 78235..87873
              /note="assembly-fragment"
misc-feature 87974..96197
              /note="assembly-fragment"
misc-feature 96298..103976
              /note="assembly-fragment clone_end:77 vector_side:left"
misc-feature 104077..110726
              /note="assembly-fragment"
misc-feature 110827..116866
              /note="assembly-fragment"
misc-feature 116867..122789
              /note="assembly-fragment"
misc-feature 122890..127665
              /note="assembly-fragment"
misc-feature 127766..132618
              /note="assembly-fragment"
misc-feature 132719..136051
              /note="assembly-fragment"
misc-feature 136152..139749
              /note="assembly-fragment"
misc-feature 139850..142810
              /note="assembly-fragment"
misc-feature 142811..145855
              /note="assembly-fragment"
misc-feature 145956..148840
              /note="assembly-fragment"
misc-feature 148941..151243
              /note="assembly-fragment"
misc-feature 151344..153454
              /note="assembly-fragment"
misc-feature 153555..156134
              /note="assembly-fragment"
misc-feature 156235..158089
              /note="assembly-fragment"
misc-feature 158190..159747
              /note="assembly-fragment"
BASE COUNT 47302 a 30397 c 29897 g 49849 t 2302 others
ORIGIN
Query Match 35.1% Score 39; DB 2; Length 159747;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
60 GGAGTATGTAACCTCCTGGTCTCTGTGTGCTGAG 98
|||||
DB 74932 GGAGTATGTAACCTCCTGGTCTCTGTGTGCTGAG 74894

RESULT 15
AC034249 162740 bp DNA linear HTG 31-AUG-2001
LOCUS Homo sapiens chromosome 5 clone RP11-427C17, WORKING DRAFT
DEFINITION
SEQUENCE, 6 ordered pieces.
AC034249
AC034249.3 GI:15383785
KEYWORDS
HTG; HTGS_PHASE2; HTGS_DRAFT; HTGS_ACTIVEFIN.
SOURCE
Homo sapiens.
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE
1 (bases 1 to 162740)
AUTHORS
DOE Joint Genome Institute.
TITLE
Sequencing of Human Chromosome 5
JOURNAL
Unpublished
REFERENCE
2 (bases 1 to 162740)
```

```
AUTHORS
TITLE
JOURNAL
COMMENT
DOE Joint Genome Institute.
Direct Submission
Submitted (05-APR-2000) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
On Aug 31, 2001 this sequence version replaced gi:9211234.
-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov
-----
Project Information
Center Project Name: 570398
Center clone name: RPCI-11_427C17
-----
Summary Statistics
Consensus quality: 158733 bases at least Q40
Consensus quality: 161469 bases at least Q30
Consensus quality: 162087 bases at least Q20
Estimated insert size: 160000; pulse field gel estimation
Estimated insert size: 162240; sum-of-contigs estimation
Quality coverage: 9.16 in Q20 bases; pulse field gel estimation
Quality coverage: 9.16 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. Gaps between the contigs
* are represented as runs of N. The order of the pieces
* is believed to be correct as given, however the sizes
* of the gaps between them are based on estimates that have
* provided by the submitter.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
1 85871: contig of 85871 bp in length
85872 96765: contig of 10794 bp in length
85972 96765: contig of 10794 bp in length
96766 96865: gap of unknown length
96866 103302: contig of 6437 bp in length
103303 103402: gap of unknown length
103403 111971: contig of 8569 bp in length
111972 112071: gap of unknown length
112072 129034: contig of 16963 bp in length
129035 129135: gap of unknown length
129135 162740: contig of 33606 bp in length.
*
FEATURES
Location/Qualifiers
source
1..162740
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="RP11-427C17"
/clone_lib="RPCI human BAC library 11"
BASE COUNT 48722 a 31435 c 31563 g 50520 t 500 others
ORIGIN
Query Match 35.1% Score 39; DB 2; Length 162740;
Best Local Similarity 100.0%; Pred. No. 1.6e-11;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
61 GAGTATGTAACCTCCTGGTCTCTGTGTGCTGAGT 99
|||||
DB 3599 GAGTATGTAACCTCCTGGTCTCTGTGTGCTGAGT 3637

Search completed: April 25, 2003, 00:30:53
Job time : 719.885 secs
```


GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:19:38 ; Search time 51.0118 Seconds
(without alignments)
4900.271 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggtgagatcttttgcctt.....gcctgagtgctgcttact 111

Scoring table: OLIGO-NUC
Gapop 60.0 , Gapext 60.0

Searched: 2185239 segs, 1125999159 residues

Read size: 0
Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Database :

N_Geneseq_101002:*

- 1: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1980.DAT:*
- 2: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1981.DAT:*
- 3: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1982.DAT:*
- 4: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1983.DAT:*
- 5: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1984.DAT:*
- 6: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1985.DAT:*
- 7: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1986.DAT:*
- 8: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1987.DAT:*
- 9: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1988.DAT:*
- 10: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1989.DAT:*
- 11: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1990.DAT:*
- 12: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1991.DAT:*
- 13: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1992.DAT:*
- 14: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1993.DAT:*
- 15: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1994.DAT:*
- 16: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1995.DAT:*
- 17: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1996.DAT:*
- 18: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1997.DAT:*
- 19: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1998.DAT:*
- 20: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA1999.DAT:*
- 21: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2000.DAT:*
- 22: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001A.DAT:*
- 23: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
- 24: /SIDS2/gcgcdata/geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	447	21 AAC03794	Human secreted pro
2	111	100.0	447	21 AAZ42680	Human 5' EST isola
3	24	21.6	660	23 AAS73441	DNA encoding novel
4	20	18.0	2526	23 AAS66674	DNA encoding novel
5	18	16.2	431	24 ABR24383	DNA encoding human
6	18	16.2	431	24 ABR24390	DNA encoding human
7	18	16.2	570	22 ABR63453	Human foetal liver
8	18	16.2	570	22 ABR30652	Probe #9118 for ge
9	18	16.2	570	22 AAK11985	Human brain expres

C 10	18	16.2	570	22 AAK37688	Human bone marrow
C 11	18	16.2	570	22 AAI18447	Probe #8380 for ge
C 12	18	16.2	570	22 AAI43563	Probe #12249 used
C 13	18	16.2	570	24 ABA11680	Human genome-deliv
C 14	18	16.2	827	23 AAS87115	DNA encoding novel
C 15	18	16.2	342	23 AAS87115	DNA encoding novel
C 16	17	15.3	266	20 AAV89709	EST clone CT857.
C 17	17	15.3	531	23 ABR43012	Genomic sequence #
C 18	17	15.3	609	24 ABR80793	Bacillus clausii g
C 19	17	15.3	828	22 AAI94407	Human neuroblastom
C 20	17	15.3	1047	24 AAS62681	CDNA sequence #468
C 21	17	15.3	1363	9 AAN70128	Novel DNA encoding
C 22	17	15.3	1546	22 AAK82125	Human immune/haema
C 23	17	15.3	1982	21 AAK68089	Human secreted pro
C 24	17	15.3	6928	22 ABA21109	Human nervous syst
C 25	17	15.3	21340	23 ABL12924	Drosophila melanog
C 26	17	15.3	62909	22 AAF28545	Genomic fragment #
C 27	17	15.3	172637	24 ABR83124	Human voltage-actl
C 28	17	15.3	495269	24 ABO67195	Listeria innocua c
C 29	17	15.3	1503900	22 AAK95240	Human neuroguilin-1
C 30	17	15.3	1503900	22 AAK96733	Human neuroguilin-1
C 31	17	15.3	3011208	24 ABO69245	Listeria innocua D
C 32	16	14.4	79	22 AAC89232	Human brain T calc
C 33	16	14.4	250	16 AAT22213	Human gene signatu
C 34	16	14.4	260	24 ABL16202	Human GREX polynuc
C 35	16	14.4	281	24 ABL67351	Thyroid cancer rel
C 36	16	14.4	302	24 ABR78873	Human ORF3820 CDNA
C 37	16	14.4	347	22 ABR60208	Human cancer relat
C 38	16	14.4	348	22 AAF65225	Novel human polynu
C 39	16	14.4	360	22 AAF57925	Human immune/haema
C 40	16	14.4	377	24 ABL38444	Human colon tumour
C 41	16	14.4	407	23 ABL36584	Human prostate exp
C 42	16	14.4	414	22 AAI82407	Human polynucleoti
C 43	16	14.4	415	23 ABO7896	Human prostate exp
C 44	16	14.4	432	23 ABL36813	Human prostate exp
C 45	16	14.4	434	22 AAI79941	Human polynucleoti

ALIGNMENTS

RESULT 1
AAC03794 standard; CDNA; 447 BP.
ID AAC03794;
AC AAC03794;
XX 06-OCT-2000 (first entry)
DT
XX Human secreted protein 5' EST, SEQ ID NO: 3792.
DE
XX Human; 5' EST; expressed sequence tag; secreted protein; CDNA isolation;
KW gene therapy; chromosome mapping; ss.
XX
XX Homo sapiens.
OS
XX
XX PN EPI033401-A2.
XX PD 06-SEP-2000.
XX PF 21-FEB-2000; 2000EP-0200610.
XX PR 26-FEB-1999; 99US-0122487.
XX (GEST) GENSET.
PA Dumas Milne Edwards J, Duclert A, Giordano J;
PI Dumas Milne Edwards J, Duclert A, Giordano J;
XX MPI: 2000-500381/45.
XX P-PSDB; AAG03788.
XX
XX New nucleic acid that is a 5' expressed sequence tag (5' EST) for
XX obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for

diagnostic, forensic, gene therapy and chromosome mapping procedures -
Claim 1: SEQ ID 3792; 71bp + CD-ROM; English.

The present sequence is one of a large number of 5' ESTs derived from cDNAs encoding secreted proteins. An ORF has been identified within the sequence. The 5' ESTs were prepared from total human RNAs or poly(A) RNAs derived from 30 different tissues. EST sequences usually correspond mainly to the 3' untranslated region (UTR) of the mRNA because they are often obtained from oligo-dT primed cDNA libraries. Such ESTs are not well suited for isolating cDNA sequences derived from the 5' ends of cDNAs and even in those cases where longer cDNA sequences have been obtained, the full 5' UTR is rarely included. 5' ESTs are derived from mRNAs with intact 5' ends and can therefore be used to obtain full length cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic, gene therapy and chromosome mapping procedures. They are used to obtain upstream regulatory sequences and to design expression and secretion vectors.

Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;

Query Match 100.0%; Score 111; DB 21; Length 447;
Best Local Similarity 100.0%; Pred. No. 3e-46; Mismatches 0; Gaps 0;
Matches 111; Conservative 0; Indels 0;

QY 1 ATGGGTGATCTTTTGCCTGAGATTCCTTTTCATCTTTGACGAGACTTGGGGCCG 60
DB 51 ATGGGTGATCTTTTGCCTGAGATTCCTTTTCATCTTTGACGAGACTTGGGGCCG 110
QY 61 GAGTATGTAACACTCCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTACT 111
DB 111 GAGTATGTAACACTCCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTACT 161

RESULT 2

AA242680 standard; cDNA; 447 BP.

AA242680;

01-FEB-2000 (first entry)

Human 5' EST isolated from a cDNA library SEQ ID NO:439.

Human: 5' EST; expressed sequence tag; secreted protein; diagnosis;
gene therapy; chromosome mapping; upstream regulatory sequence;
forensic; location; development; protein synthesis; stability;
regulation; identification; ss.

Homo sapiens.

WO9953051-A2.

21-OCT-1999.

09-APR-1999; 99WO-IB00712.

09-APR-1998; 98US-0057719.

28-APR-1998; 98US-0069047.

(GEST) GENSET.

Dumas Milne Edwards J, Duclert A, Giordano J;

WPI; 2000-038446/03.

P-PSDB; AAY65066.

Novel secreted protein 5' expressed sequence tag sequences used in
diagnostic, forensic, gene therapy, and chromosome mapping procedures

Claim 1: Page 402; 837pp; English.
AA242685 to AA243075 represent novel 5' expressed sequence tag (EST)

sequences, corresponding to human secreted proteins. AAY64651 to
AA16538 represent the EST-related proteins corresponding to AA242265 to
AA43052. The 5' ESTs can be used for producing secreted human gene
products. They can be used to identify and isolate 5' untranslated
regions (UTRs) and upstream regulatory regions which control the
location, development stage, rate, and quantity of protein synthesis, as
well as stability of mRNA. The ESTs are also useful as probes for
chromosome mapping, and to obtain full length cDNA clones. The ESTs can
also be used in forensic procedures to identify individuals, or in
diagnostic procedures to identify individuals having genetic diseases
resulting from abnormal gene expression. The products may also be used in
gene therapy protocols. The nucleic acids encoding signal peptides can be
used for directing extracellular secretion of a polypeptide or the
insertion of a polypeptide into a membrane, or importing a polypeptide
into a cell. The proteins encoded by the EST sequences may be useful in
treating a variety of human conditions. Secreted proteins have
therapeutic value, and the identification of new secreted proteins is
valuable. AA242249 to AA242264 and AAY64644 to AAY64650 represent
sequences used in the exemplification of the present invention.

Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;

Query Match 100.0%; Score 111; DB 21; Length 447;
Best Local Similarity 100.0%; Pred. No. 3e-46; Mismatches 0; Gaps 0;
Matches 111; Conservative 0; Indels 0;

QY 1 ATGGGTGATCTTTTGCCTGAGATTCCTTTTCATCTTTGACGAGACTTGGGGCCG 60
DB 51 ATGGGTGATCTTTTGCCTGAGATTCCTTTTCATCTTTGACGAGACTTGGGGCCG 110
QY 61 GAGTATGTAACACTCCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTACT 111
DB 111 GAGTATGTAACACTCCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTACT 161

RESULT 3

AA573441/C standard; cDNA; 660 BP.

AA573441;

13-FEB-2002 (first entry)

DNA encoding novel human diagnostic protein #9245.

Human: chromosome mapping; gene mapping; gene therapy; forensic;
food supplement; medical imaging; diagnostic; genetic disorder; ss.
Homo sapiens.

WO200175067-A2.

11-OCT-2001.

30-MAR-2001; 2001WO-US08631.

31-MAR-2000; 2000US-0540217.

23-AUG-2000; 2000US-0649167.

(HYSE-) HYSED INC.

Dzmanac RT, Liu C, Tang YT;

WPI; 2001-639362/73.

P-PSDB; ABG09254.

New isolated polynucleotide and encoded polypeptides, useful in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits and to assess
biodiversity

Claim 1: SEQ ID NO 9245; 103pp; English.

CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

Sequence 660 BP: 149 A; 127 C; 148 G; 236 T; 0 other;

Query Match 21.6%; Score 24; DB 23; Length 660;
Best Local Similarity 100.0%; Pred. No. 0.017;
Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 69 AAAACTCTGGGCTCTGTGTGTG 92
|||||
Db 656 AAAACTCTGGGCTCTGTGTGTG 633

RESULT 4
AAS66674
ID AAS66674 standard; cDNA; 2526 BP.

XX AAS66674;

DT 13-FEB-2002 (first entry)

DE DNA encoding novel human diagnostic protein #2478.

XX Human; chromosome mapping; gene mapping; gene therapy; forensic;
KM food supplement; medical imaging; diagnostic; genetic disorder; ss.

XX Homo sapiens.

OS WO200175067-A2.

XX 11-OCT-2001.

PF 30-MAR-2001; 2001WO-US08631.

PR 31-MAR-2000; 2000US-0540217.

PR 23-AUG-2000; 2000US-0649167.

XX (HYSE-) HYSEQ INC.

PI Dmanac RT, Liu C, Tang YT;

DR WPI; 2001-639362/73.

DR P-PSDB; ABG02487.

XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity

XX Claim 1; SEQ ID No 2478; 103pp; English.

XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,

CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

Sequence 2526 BP: 871 A; 536 C; 541 G; 578 T; 0 other;

Query Match 18.0%; Score 20; DB 23; Length 2526;
Best Local Similarity 100.0%; Pred. No. 1.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 32 TTTCATCTTTCGAGGACTT 51
|||||
Db 2377 TTTCATCTTTCGAGGACTT 2396

RESULT 5
ABR24383
ID ABR24383 standard; cDNA; 431 BP.

XX ABR24383;

DT 09-APR-2002 (first entry)

DE DNA encoding human lung cancer protein, Seq ID No 80.

XX Human; lung cancer; cytostatic; vaccine; gene; ss.

XX Homo sapiens.

PN WO200192525-A2.

PD 06-DEC-2001.

PF 25-MAY-2001; 2001WO-US17066.

PR 26-MAY-2000; 2000US-207485P.

PR 06-SEP-2000; 2000US-230475P.

XX (CORI-) CORIXA CORP.

PI Harlocker SL, Wang T, Bangur CS, Klee JT, Switzer A;

DR WPI; 2002-122068/16.

XX New lung tumour polypeptides and polynucleotides, useful in
PT pharmaceutical compositions, such as vaccines, for treating or
PT preventing lung cancer, or as probes or primers for nucleic acid
PT hybridisation

XX Claim 9; Page 172; 179pp; English.

XX The invention relates to novel human lung cancer polynucleotide (I)
CC and polypeptides (II). (I) and (II) are useful in pharmaceutical
CC compositions, such as vaccines, for the diagnosis and treatment of lung
CC cancer. The polynucleotides are also useful as probes or primers for
CC nucleic acid hybridisation. ABR24314-ABR24397 represent human lung
CC cancer coding sequences of the invention.

XX Sequence 431 BP; 108 A; 97 C; 123 G; 101 T; 2 other;
SQ
Query Match 16.2%; Score 18; DB 24; Length 431;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 91 TGCCTGAGTGGCTGCTCT 108
|||||
Db 18 TGCCTGAGTGGCTGCTCT 35

RESULT 6
ABK24390
ID ABK24390 standard; CDNA; 431 BP.
XX
AC ABK24390;
XX
DT 09-APR-2002 (first entry)

XX DNA encoding human lung cancer protein, Seq ID No 87.
XX
XX Human; lung cancer; cytostatic; vaccine; gene; ss.
XX
XX Homo sapiens.
XX
XX WO200192525-A2.
XX
XX PD 06-DEC-2001.
XX
XX PF 25-MAY-2001; 2001WO-US17066.
XX
XX PR 26-MAY-2000; 2000US-207485P.
XX
XX PR 06-SEP-2000; 2000US-230475P.
XX
XX PA (CORI-) CORIXA CORP.
XX
XX PI Harlocker SL, Wang T, Bangur CS, Klee JT, Switzer A;
XX
XX WPI: 2002-122068/16.
XX
XX DR New lung tumour polypeptides and polynucleotides, useful in
XX
XX PT pharmaceutical compositions, such as vaccines, for treating or
XX
XX PT preventing lung cancer, or as probes or primers for nucleic acid
XX
XX PT hybridisation
XX
XX PS Claim 1; Page 174; 179pp; English.

XX The invention relates to novel human lung cancer polynucleotide (I)
XX and polypeptides (II). (I) and (II) are useful in pharmaceutical
XX compositions, such as vaccines, for the diagnosis and treatment of lung
XX cancer. The polynucleotides are also useful as probes or primers for
XX nucleic acid hybridisation. ABK24314-ABK24397 represent human lung
XX cancer coding sequences of the invention.
XX
XX SQ Sequence 431 BP; 108 A; 97 C; 123 G; 101 T; 2 other;
XX
XX Query Match 16.2%; Score 18; DB 24; Length 431;
XX
XX Best Local Similarity 100.0%; Pred. No. 17;
XX
XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 91 TGCCTGAGTGGCTGCTCT 108
XX
XX |||||
XX
XX Db 18 TGCCTGAGTGGCTGCTCT 35

RESULT 7
ABA63453/c
ID ABA63453 standard; DNA; 570 BP.
XX
AC ABA63453;
XX
XX
XX DT 01-FEB-2002 (first entry)

XX
DE Human foetal liver single exon nucleic acid probe #11758.
XX
XX KW Human; foetal liver; gene expression; single exon nucleic acid probe; ss.
XX
XX OS Homo sapiens.
XX
XX PN WO200157277-A2.
XX
XX PD 09-AUG-2001.
XX
XX PF 30-JAN-2001; 2001WO-US00669.
XX
XX PR 04-FEB-2000; 2000US-0180312.
XX
XX PR 26-MAY-2000; 2000US-0207456.
XX
XX PR 30-JUN-2000; 2000US-0608408.
XX
XX PR 03-AUG-2000; 2000US-0632366.
XX
XX PR 21-SEP-2000; 2000US-0234687.
XX
XX PR 27-SEP-2000; 2000US-0236359.
XX
XX PR 04-OCT-2000; 2000GB-0024263.
XX
XX PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX DR WPI: 2001-483447/52.
XX
XX PT Human genome-derived single exon nucleic acid probes useful for
XX
XX PT analyzing gene expression in human fetal liver -
XX
XX PS Claim 1; SEQ ID NO 11758; 639pp + sequence listing; English.
XX
XX CC The invention relates to a single exon nucleic acid probe for
XX
XX CC measuring human gene expression in a sample derived from human foetal
XX
XX CC liver. The single exon nucleic acid probes may be used for predicting,
XX
XX CC measuring and displaying gene expression in samples derived from human
XX
XX CC fetal liver. The present sequence is a single exon nucleic acid
XX
XX CC probe of the invention.
XX
XX CC Note: The sequence data for this patent did not form part of the
XX
XX CC printed specification, but was obtained in electronic format directly
XX
XX CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.
XX
XX SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
XX
XX Query Match 16.2%; Score 18; DB 22; Length 570;
XX
XX Best Local Similarity 100.0%; Pred. No. 17;
XX
XX Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 85 TGTGTGCTGCTGAGTGGC 102
XX
XX |||||
XX
XX Db 505 TGTGTGCTGCTGAGTGGC 488

RESULT 8
ABA30652/c
ID ABA30652 standard; DNA; 570 BP.
XX
AC ABA30652;
XX
XX
XX DT 23-JAN-2002 (first entry)
XX
XX DE Probe #9118 for gene expression analysis in human heart cell sample.
XX
XX DE Human; gene expression; heart; microarray; vascular system; probe;
XX
XX KW cardiovascular disease; hypertension; cardiac arrhythmia.
XX
XX KW congenital heart disease; ss.
XX
XX OS Homo sapiens.
XX
XX XX
XX PN WO200157274-A2.
XX
XX PD 09-AUG-2001.

PF 30-JAN-2001; 2001WO-US00666.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488990/53.
XX
PT Single exon nucleic acid probes for analyzing gene expression in human
PT hearts -
XX
PS Claim 1; SEQ ID No 9118; 530pp; English.
XX
CC The present invention relates to single exon nucleic acid probes for
CC measuring human gene expression in a sample derived from human heart. The
CC present sequence is one such probe. The probes may be used for
CC predicting, measuring and displaying gene expression in samples derived
CC from the human heart via microarrays. By measuring gene expression, the
CC probes are useful for predicting, diagnosing, grading, staging,
CC monitoring and prognosing diseases of the human heart and vascular system
CC e.g. cardiovascular disease, hypertension, cardiac arrhythmias and
CC congenital heart disease.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
XX
Query Match 16.2%; Score 18; DB 22; Length 570;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 85 TGTGTGTCCTGAGTGGC 102
XXXXXXXXXXXXXXXXXXXX
Db 505 TGTGTGTCCTGAGTGGC 488
XX
RESULT 9
AAK1985/c
ID AAK1985 standard; DNA; 570 BP.
XX
AAK1985;
XX
DT 05-NOV-2001 (first entry)
XX
XX Human brain expressed single exon probe SEQ ID NO: 11976.
DE
XX Human brain expressed single exon probe SEQ ID NO: 11976.
XX
KW Human; brain expressed exon; gene expression analysis; probe;
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
KW epilepsy; cancer; ss.
XX
XX Homo sapiens.
OS
XX
PN WO200157275-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00667.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX

PR 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
PA
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-483446/52.
XX
DR Single exon nucleic acid probes for analyzing gene expression in human
XX brains -
PT
PT
PS Example 4; SEQ ID NO: 11976; 650pp + Sequence Listing; English.
XX
XX The present invention provides a number of single exon nucleic acid
XX probes which are derived from genomic sequences expressed in the human
XX brain. They can be used to measure gene expression in brain cell samples,
XX which may enable the diagnosis and improved treatment of nervous system
XX diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
XX epilepsy and cancers. The present sequence is one of the probes of the
XX invention.
XX
SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
XX
Query Match 16.2%; Score 18; DB 22; Length 570;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
QY 85 TGTGTGTCCTGAGTGGC 102
XXXXXXXXXXXXXXXXXXXX
Db 505 TGTGTGTCCTGAGTGGC 488
XX
RESULT 10
AAK37688/c
ID AAK37688 standard; DNA; 570 BP.
XX
XX AAK37688;
XX
DT 06-NOV-2001 (first entry)
XX
XX Human bone marrow expressed single exon probe SEQ ID NO: 12245.
DE
XX Human bone marrow expressed single exon probe SEQ ID NO: 12245.
XX
KW Human; bone marrow expressed exon; gene expression analysis; probe;
KW microarray; cancer; leukemia; lymphoma; myeloma; ss.
XX
XX Homo sapiens.
OS
XX
PN WO200157276-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US00668.
XX
PR 04-FEB-2000; 2000US-0180312.
PR 26-MAY-2000; 2000US-0207456.
PR 30-JUN-2000; 2000US-0608408.
PR 03-AUG-2000; 2000US-0632366.
PR 21-SEP-2000; 2000US-0234687.
PR 27-SEP-2000; 2000US-0236359.
PR 04-OCT-2000; 2000GB-0024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
PA
PI Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488990/53.
XX
DR Human genome-derived single exon nucleic acid probes useful for
XX analyzing gene expression in human bone marrow -
PT
PT Example 4; SEQ ID NO: 12245; 658pp + Sequence Listing; English.
XX
XX The present invention provides a number of single exon nucleic acid
XX

CC probes which are derived from genomic sequences expressed in the human
CC bone marrow. They can be used to measure gene expression in bone marrow
CC samples, which may enable the improved diagnosis and treatment of cancers
CC such as lymphoma, leukaemia and myeloma. The present sequence is one of
CC the probes of the invention.

SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 16.2%; Score 18; DB 22; Length 570;

Best Local Similarity 100.0%; Pred. No. 17;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 85 TGTGTGTCCTGAGTGGC 102

Db 505 TGTGTGTCCTGAGTGGC 488

RESULT 11

AA118447/c

AA118447 standard; DNA; 570 BP.

AA118447;

DT 12-OCT-2001 (first entry)

DE Probe #8380 for gene expression analysis in human cervical cell sample.

XX Probe: human; microarray; gene expression; cervical epithelial cell;

KW cervical cancer; ss.

XX Homo sapiens.

OS WO200157278-A2.

PN 09-AUG-2001.

PD 30-JAN-2001; 2001WO-US00670.

XX 04-FEB-2000; 2000US-0180312.

XX 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

PA Penn SG, Hanzel DK, Chen W, Rank DR;

DR WPI; 2001-488901/53.

XX Human genome-derived single exon nucleic acid probes useful for

PT analyzing gene expression in human cervical epithelial cells -

XX Claim 25; SEQ ID No 8380; 487bp; English.

XX The present invention relates to human single exon nucleic acid probes

CC (SENP). The present sequence is one such probe. The SENPs are derived

CC from human HeLa cells. The SENPs can be used to produce a single exon

CC microarray, which can be used for measuring human gene expression in a

CC sample derived from human cervical epithelial cells. By measuring gene

CC expression, the probes are therefore useful in grading and/or staging

CC of diseases of the cervix, notably cervical cancer.

CC Note: The sequence data for this patent did not form part of the printed

CC specification, but was obtained in electronic format directly from WIPO

CC at ftp.wipo.int/pub/published_pct_sequences.

XX Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

SQ Query Match 16.2%; Score 18; DB 22; Length 570;

Best Local Similarity 100.0%; Pred. No. 17;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 85 TGTGTGTCCTGAGTGGC 102

Db 505 TGTGTGTCCTGAGTGGC 488

RESULT 12

AA143563/c

ID AA143563 standard; DNA; 570 BP.

XX AA143563;

DT 17-OCT-2001 (first entry)

DE Probe #12249 used to measure gene expression in human placenta sample.

XX Probe: microarray; human; placenta; antenatal diagnosis;

KW genetic disorder; ss.

XX Homo sapiens.

OS WO200157272-A2.

PN 09-AUG-2001.

PD 30-JAN-2001; 2001WO-US00663.

XX 04-FEB-2000; 2000US-0180312.

XX 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

PA Penn SG, Hanzel DK, Chen W, Rank DR;

DR WPI; 2001-48897/53.

XX Human genome-derived single exon nucleic acid probes useful for

PT analyzing gene expression in human placenta -

XX Claim 25; SEQ ID No 12249; 654bp; English.

XX The present invention relates to single exon nucleic acid probes (SENP).

CC The present sequence is one such probe. The probes are useful for

CC producing a microarray for predicting, measuring and displaying gene

CC expression in samples derived from human placenta. The probes are useful

CC for antenatal diagnosis of human genetic disorders.

XX Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

SQ Query Match 16.2%; Score 18; DB 22; Length 570;

Best Local Similarity 100.0%; Pred. No. 17;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 85 TGTGTGTCCTGAGTGGC 102

Db 505 TGTGTGTCCTGAGTGGC 488

RESULT 13

ABSI1680/c

ID ABSI1680 standard; DNA; 570 BP.

XX ABSI1680;

DT 19-AUG-2002 (first entry)

DE Human genome-derived single exon probe from lung SEQ ID No 11671.

KW Human: ds; single exon probe; asthma; lung cancer; COPD; ILD;
 KW chronic obstructive pulmonary disease; interstitial lung disease;
 KW familial idiopathic pulmonary fibrosis; neurofibromatosis;
 KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
 KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemorrhoidosis;
 KW pulmonary histiocytosis; lymphangioleiomyomatosis; Kargener syndrome;
 KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
 KW primary ciliary dyskinesia; pulmonary hypertension;
 KW hyaline membrane disease.
 XX
 OS Homo sapiens.
 PN WO200186003-A2.
 XX
 PD 15-NOV-2001.
 XX
 PF 30-JAN-2001; 2001WO-US000665.
 XX
 PR 04-FEB-2000; 2000US-180312P.
 PR 26-MAY-2000; 2000US-207456P.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0632366.
 PR 21-SEP-2000; 2000US-234687P.
 PR 27-SEP-2000; 2000US-236359P.
 PR 04-OCT-2000; 2000GB-0024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 XX
 PI Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 DR WPI; 2002-114183/15.
 XX
 PT Spatially-addressable set of single exon nucleic acid probes, used to
 PT measure gene expression in human lung samples -
 XX
 PS Claim 1; SEQ ID No 11671; 634pp; English.
 XX
 CC The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human lung comprising single exon nucleic acid probes having one of
 CC 12614 nucleic acid sequences mentioned in the specification, or their
 CC complements or the 12387 open reading frames derived from the 12614
 CC probes. Also included are a microarray comprising the novel set of
 CC probes; the novel set of probes which hybridise at high stringency to a
 CC nucleic acid expressed in the human lung; measuring gene expression in a
 CC sample derived from human lung, comprising (a) contacting the array with
 CC a collection of detectably labeled nucleic acids derived from human lung
 CC mRNA, and (b) measuring the label detectably bound to each probe of
 CC the array; identifying exons in a eukaryotic genome, comprising
 CC (a) algorithmically predicting at least one exon from genomic sequences
 CC of the eukaryote; and (b) detecting specific hybridisation of detectably
 CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
 CC having a fragment identical to the predicted exon, the probe is included
 CC in the above mentioned microarray, assigning exons to a single gene,
 CC comprising (a) identifying exons from genomic sequence by the method
 CC above and (b) measuring the expression of each of the exons in several
 CC tissues and/or cell types using hybridisation to a single exon
 CC microarrays having a probe with the exon, where a common pattern of
 CC expression of the exons in the tissues and/or cell types indicates that
 CC the exons should be assigned to a single gene; a peptide comprising one
 CC of 12011 sequences, mentioned in the specification, or encoded by the
 CC probes/open reading frames (ORF). The probes are used for gene
 CC expression analysis, and for identifying exons in a gene, particularly
 CC using human lung derived mRNA and for the study of lung diseases
 CC such as asthma, lung cancer, chronic obstructive pulmonary disease
 CC (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
 CC fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
 CC Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
 CC haemorrhoidosis, pulmonary histiocytosis, lymphangioleiomyomatosis,
 CC pulmonary alveolar proteinosis, Kargener syndrome, fibrocystic
 CC pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
 CC and hyaline membrane disease. The present sequence is a single exon
 CC probe of the invention.

CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX
 S0 Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
 Query Match 16.2%; Score 18; DB 24; Length 570;
 Best Local Similarity 100.0%; Pred. No. 17;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 0Y 85 TGTGTGTCCTGAGTGC 102
 ||||||||||||||||
 Db 505 TGTGTGTCCTGAGTGC 488
 RESULT 14
 AAS87115
 ID AAS87115 standard; cDNA; 827 BP.
 XX
 AC AAS87115;
 XX
 DT 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #22919.
 XX
 KW Human: chromosome mapping; gene mapping; gene therapy; forensic;
 KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 OS Homo sapiens.
 XX
 PN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 PF 30-MAR-2001; 2001WO-US008631.
 XX
 PR 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 PA (HSE-) HSEQ INC.
 XX
 PI Drmanac RT, Liu C, Tang YT;
 XX
 DR WPI; 2001-639362/73.
 XX
 PT P-PSDB; ABG22928.
 XX
 PS Claim 1; SEQ ID No 22919; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC a food supplement. (II) and its binding partners are useful in medical
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed

CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 827 BP; 198 A; 202 C; 249 G; 178 T; 0 other;
Query Match 16.2%; Score 18; DB 23; Length 827;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 91 TGCCTGAGTGCTGCTCT 108
|||||
Db 344 TGCCTGAGTGCTGCTCT 361
RESULT 15
AAS87118/c
ID AAS87118 standard; cDNA; 3342 BP.
XX
PC AAS87118;
XX
13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #22922.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
PA (HYSE-) HYSEQ INC.
XX
PI Drmanac RT, Liu C, Tang YT;
XX
DR WPI: 2001-639362/73.
DR P-PSDB; ABG22931.
XX
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
diagnostics, forensics, gene mapping, identification of mutations
responsible for genetic disorders or other traits and to assess
biodiversity
XX
PS Claim 1; SEQ ID No 22922; 103pp; English.
XX
XX
CC The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS64197-AAS94564 represent novel human
CC diagnostic coding sequences. AAS64197-AAS94564 represent novel human
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.

XX
SQ Sequence 3342 BP; 898 A; 796 C; 867 G; 781 T; 0 other;
Query Match 16.2%; Score 18; DB 23; Length 3342;
Best Local Similarity 100.0%; Pred. No. 17;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 91 TGCCTGAGTGCTGCTCT 108
|||||
Db 113 TGCCTGAGTGCTGCTCT 96

Search completed: April 25, 2003, 00:00:24
Job time : 56.0118 secs

GenCore version 5.1.4.p5_4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:49:04 ; Search time 373.284 seconds
(without alignments)
4815.901 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggtggtatcttcttgcctt.....gcctgagtgcgtcttact 111

Scoring table: OLIGO/MUC
Gapop 60.0 , Gapext 60.0

Searched: 16154066 seqs, 8097743376 residues

Database size : 0
Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0
Maximum DB seq length: 200000000

Post-processing: Listing first 45 summaries

Database :
EST:*
1: em_estbda:*
2: em_esthum:*
3: em_estin:*
4: em_estnu:*
5: em_estrov:*
6: em_estrpl:*
7: em_estro:*
8: em_hlc:*
9: gb_est1:*
10: gb_est2:*
11: gb_hlc:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: gb_gss:*
18: em_gss_hum:*
19: em_gss_iny:*
20: em_gss_pln:*
21: em_gss_vrt:*
22: em_gss_fun:*
23: em_gss_man:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rtd:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	470	AQ770688	AQ770688 HS_5368_B
2	45	40.5	525	AQ165256	AQ165256 HS_3025_B
3	38	34.2	628	AQ237815	AQ237815 RPT11-70
4	35	31.5	410	AQ442274	AQ442274 HS_5137_A
5	35	31.5	453	AQ437684	AQ437684 HS_5137_A
6	35	31.5	633	AQ390599	AQ390599 CITR1-EI

c	7	34	30.6	529	17	AQ881246	AQ881246 HS_5137_B
c	8	33	29.7	482	17	AQ320567	AQ320567 RPT11-99
c	9	33	29.7	563	17	AQ420187	AQ420187 RPT11-1
c	10	33	29.7	723	17	AQ386439	AQ386439 RPT11-15
c	11	31	27.9	553	17	AQ521751	AQ521751 RPT11-1
c	12	31	27.9	695	17	AG179297	AG179297 Pan trogl
c	13	30	27.0	360	17	AQ207172	AQ207172 HS_3239_B
c	14	30	27.0	399	17	AQ115544	AQ115544 RPT11-57
c	15	30	27.0	435	17	AQ116061	AQ116061 RPT11-57
c	16	30	27.0	551	17	AQ569689	AQ569689 HS_5333_B
c	17	29	26.1	514	17	AQ003326	AQ003326 RPT11-25
c	18	29	26.1	615	17	AG161224	AG161224 Pan trogl
c	19	28	25.2	363	17	AQ120796	AQ120796 HS_3076_A
c	20	28	25.2	376	17	AQ548294	AQ548294 RPT11-4
c	21	28	25.2	401	17	AQ568089	AQ568089 CITR1-EI
c	22	28	25.2	419	17	AQ36209	AQ36209 HS_5049_B
c	23	28	25.2	530	17	AQ193128	AQ193128 HS_3060_B
c	24	28	25.2	653	17	AG160919	AG160919 Pan trogl
c	25	27	24.3	553	17	AQ238365	AQ238365 RPT11-63
c	26	27	24.3	606	17	AQ350708	AQ350708 RPT11-11
c	27	26	23.4	478	17	B75615	B75615 RPT11-11L1
c	28	26	23.4	653	17	AG143347	AG143347 Pan trogl
c	29	26	23.4	690	17	AG11263	AG11263 Pan trogl
c	30	25	22.5	411	17	AQ715895	AQ715895 HS_5430_B
c	31	25	22.5	427	17	AQ697116	AQ697116 HS_5528_A
c	32	25	22.5	456	17	AQ707169	AQ707169 HS_5564_A
c	33	25	22.5	468	17	AQ533287	AQ533287 RPT11-3
c	34	25	22.5	519	17	AQ519549	AQ519549 HS_5168_A
c	35	25	22.5	520	17	AQ375479	AQ375479 RPT11-11
c	36	25	22.5	552	17	AQ378145	AQ378145 RPT11-15
c	37	25	22.5	563	17	AQ379371	AQ379371 RPT11-16
c	38	25	22.5	583	17	AG070759	AG070759 Pan trogl
c	39	25	22.5	609	17	AG155327	AG155327 Pan trogl
c	40	25	22.5	747	17	AG170366	AG170366 Pan trogl
c	41	24	21.6	413	17	AQ26658	AQ26658 RPT11-73
c	42	24	21.6	425	17	AQ683450	AQ683450 HS_5432_B
c	43	24	21.6	471	17	AQ677950	AQ677950 HS_5331_B
c	44	24	21.6	506	17	AQ284282	AQ284282 RPT11-79
c	45	24	21.6	541	17	AQ686074	AQ686074 HS_5537_B

ALIGNMENTS

RESULT 1
LOCUS AQ770688 470 bp DNA linear GSS 28-JUN-1999
DEFINITION HS_5368_B2_C08-SP66 RPT11-11 Human Male BAC Library Homo sapiens
ACCESSION genomic clone Plate-944 Col-16 Row-F, DNA sequence.
VERSION AQ770688.1 GI:5648804
KEYWORDS GSS.

ORGANISM

human.

REFERENCE
1 (bases 1 to 470)
Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

AUTHORS

Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.

Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

JOURNAL

CONTACT: Mahairas GG, Wallace JC, Hood L

COMMENT

High Throughput Sequencing Center

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPT11-11. For BAC
library availability, please contact Pieter de Jong

(pieterdejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm) or from Research Genetics (info@resgen.com). BAC end Web Server: <http://www.husc.washington.edu>

Plate: 944 row: F column: 16
Seq primer: SP6
Class: BAC ends

High quality sequence stop: 470.

FEATURES

Source

1..470

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate=944 Col=16 Row=F"

/clone_lib="RPCI-11 Human Male BAC Library"

/sex="male"

/note="Vector: pBAC3.6; Site1: EcoRI; Site2: EcoRI; Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBAC3.6 vector at EcoRI sites"

E COUNT

83 a 112 c 131 g 141 t 3 others

ORIGIN

Query Match

Best Local Similarity 100.0%; Score 111; DB 17; Length 470;

Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

1 ATGGGAGATCTTTGCGCTGAGATCTTTTATCTTTGACGAGCTTGCGGCCG 60

Db

103 ATGGGAGATCTTTGCGCTGAGATCTTTTATCTTTGACGAGCTTGCGGCCG 162

QY

61 GAGTATGTAACCTCTGCGCTCTGTGTGCTGAGTGGCTGCTACT 111

Db

163 GAGTATGTAACCTCTGCGCTCTGTGTGCTGAGTGGCTGCTACT 213

RESULT 2

LOCUS

A0165256 525 bp DNA linear GSS 16-OCT-1998

DEFINITION

HS.3025.B2.G06.T7 CIT Approved Human Genomic Sperm Library D Homo

ACCESSION

A0165256 sapiens genomic clone Plate=3025 Col=12 Row=N, DNA sequence.

VERSION

A0165256.1 GI:3563451

KEYWORDS

GSS.

SOURCE

human.

ORGANISM

Homo sapiens

REFERENCE

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 525)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D., and
Hood,L.

TITLE

Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome

JOURNAL

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

MEDLINE

99380589

COMMENT

Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Sequence Tagged Connector
Plate: 3025 row: N column: 12
Class: BAC ends
High quality sequence stop: 525.

FEATURES

Source

1..525

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate=3025 Col=12 Row=N"

/clone_lib="CIT Approved Human Genomic Sperm Library D"

/sex="male"

/note="Organ: sperm; Vector: pBelosBAC11; BAC Clones in
E-Coli DH10B"

BASE COUNT 102 a 139 c 137 g 143 t 4 others

ORIGIN

Query Match

Best Local Similarity 100.0%; Score 45; DB 17; Length 525;

Matches 45; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

34 TCATCTTTCGAGGACCTTCGGCGCGAGATGTAACCTCCG 78

Db

99 TCATCTTTCGAGGACCTTCGGCGCGAGATGTAACCTCCG 143

RESULT 3

LOCUS

A0237815

DEFINITION

RPC111-70H4.TK RPCI-11 Homo sapiens genomic clone RPCI-11-70H4, DNA

ACCESSION

A0237815

VERSION

A0237815.1 GI:3670106

KEYWORDS

GSS.

SOURCE

human.

ORGANISM

Homo sapiens

REFERENCE

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS

Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 628)
Adams,M.D., Rounsley,S.D., Zhao,S., Bass,S., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., Wible,C., de Jong,P. and Venter,J.C.
Use of human BAC End Sequences for Sequence-Ready Map Building
Unpublished (1998)

TITLE

Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208

JOURNAL

Email: maddams@tigr.org
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact pieter de Jong
(pieterdejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from
Research Genetics (info@resgen.com). BAC end search page:
http://www.tigr.org/cdb/humgen/bac_end_search.html

COMMENT

Seq primer: T7
Class: BAC ends.

FEATURES

Source

1..628

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="RPCI-11-70H4"

/clone_lib="RPCI-11"

/sex="male"

/cell_type="Lymphocytes"

/note="Vector: pBAC3.6; Site1: EcoRI; Site2: EcoRI;

RPC111 Human Male BAC Library"

BASE COUNT 125 a 150 c 164 g 189 t

ORIGIN

Query Match

Best Local Similarity 100.0%; Score 38; DB 17; Length 628;

Matches 38; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

61 GAGTATGTAACCTCTGCGCTCTGTGTGCTGAG 98

Db

157 GAGTATGTAACCTCTGCGCTCTGTGTGCTGAG 194

RESULT 4

LOCUS

A0442274

410 bp DNA linear GSS 31-MAR-1999

DEFINITION HS_5137_A1-F12.SP6E RPCI-11 Human Male BAC Library Homo sapiens
 accession A0442274
 version A0442274.1 GI:453613
 keywords GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 410)
 Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
 Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
 Hood,L.
 TITLE Sequence-tagged connectors: A sequence approach to mapping and
 scanning the human genome
 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
 MEDLINE 99380589
 COMMENT Contact: Mahairas GG, Wallace JC, Hood L
 High Throughput Sequencing Center
 University of Washington
 401 Queen Anne Avenue North, Seattle, WA 98109, USA
 Tel: (206) 616-3618
 Fax: (206) 616-3887
 Email: jwallace@u.washington.edu
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pieter de Jong
 (pieter@dejong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering.bac.htm)
 or from Research Genetics (info@resgen.com). BAC end Web Server:
 http://www.htsc.washington.edu
 Seq primer: SP6
 Plate: 713 row: K column: 23
 Class: BAC ends
 High quality sequence stop: 410.
 FEATURES
 Location/Qualifiers
 source 1..410
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="Plate=713 Col=23 Row=K"
 /clone_lib="RPCI-11 Human Male BAC Library"
 /sex="male"
 /note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
 Male blood DNA was isolated from one randomly chosen donor
 and partially digested with a combination of EcoRI and
 EcoRI Methylase. Size selected DNA was cloned into the
 pBACe3.6 vector at EcoRI sites"
 BASE COUNT 80 a 111 c 111 g 107 t 1 others
 ORIGIN
 Query Match 31.5%; Score 35; DB 17; Length 410;
 Best Local Similarity 100.0%; Pred. No. 1.2e-07;
 Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 74 TCCTGGGCTCTGTGTGCTGAGTGCTGCTCT 108
 ||||||||||||||||||||||||||||||||
 Db 160 TCCTGGGCTCTGTGTGCTGAGTGCTGCTCT 194
 RESULT 5
 LOCUS A0437684 453 bp DNA linear GSS 31-MAR-1999
 DEFINITION HS_5137_A2.SP6E RPCI-11 Human Male BAC Library Homo sapiens
 accession A0437684
 version A0437684.1 GI:4549023
 keywords GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 453)
 Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
 Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
 Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and

TITLE Hood,L.
 JOURNAL Sequence-tagged connectors: A sequence approach to mapping and
 scanning the human genome
 Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
 MEDLINE 99380589
 COMMENT Contact: Mahairas GG, Wallace JC, Hood L
 High Throughput Sequencing Center
 University of Washington
 401 Queen Anne Avenue North, Seattle, WA 98109, USA
 Tel: (206) 616-3618
 Fax: (206) 616-3887
 Email: jwallace@u.washington.edu
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pieter de Jong
 (pieter@dejong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering.bac.htm)
 or from Research Genetics (info@resgen.com). BAC end Web Server:
 http://www.htsc.washington.edu
 Seq primer: SP6
 Plate: 713 row: O column: 12
 Class: BAC ends
 High quality sequence stop: 453.
 FEATURES
 Location/Qualifiers
 source 1..453
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="Plate=713 Col=12 Row=O"
 /clone_lib="RPCI-11 Human Male BAC Library"
 /sex="male"
 /note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
 Male blood DNA was isolated from one randomly chosen donor
 and partially digested with a combination of EcoRI and
 EcoRI Methylase. Size selected DNA was cloned into the
 pBACe3.6 vector at EcoRI sites"
 BASE COUNT 84 a 127 c 117 g 124 t 1 others
 ORIGIN
 Query Match 31.5%; Score 35; DB 17; Length 453;
 Best Local Similarity 100.0%; Pred. No. 1.2e-07;
 Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 74 TCCTGGGCTCTGTGTGCTGAGTGCTGCTCT 108
 ||||||||||||||||||||||||||||||||
 Db 163 TCCTGGGCTCTGTGTGCTGAGTGCTGCTCT 197
 RESULT 6
 LOCUS A0390599/c 635 bp DNA linear GSS 06-MAR-1999
 DEFINITION CITBI-E1-2544B15.TR CITBI-E1 Homo sapiens genomic clone 2544B15,
 accession A0390599
 version A0390599.1 GI:4361622
 keywords GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 1 (bases 1 to 635)
 Zhao,S., Adams,M.D., Nierman,W., Malek,J., Shizuya,H., Simon,M. and
 Venter,J.C.
 TITLE Use of BAC End Sequences from Caltech Libraries for Sequence-Ready
 Map Building
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: CITBI-E1-2544B15.TR
 Contact: Shaying Zhao, William Nierman, Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: bheerflg.org
 Clones are available from Research Genetics (info@resgen.com). BAC

end search page:
http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html.
Seq primer: M13 Reverse
Class: BAC ends.

FEATURES
Source

Location/Qualifiers
1. 635
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="2544B15"
/clone_lib="CITBI-E1"
/sex="male"
/cell_type="sperm"
/note="Vector: pBeloBAC11; Site_1: EcoRI; site_2: EcoRI;
Caltech Human BAC Library D"
BASE COUNT 192 a 144 c 172 g 127 t
ORIGIN

Query Match 31.5%; Score 35; DB 17; Length 635;
Best Local Similarity 100.0%; Pred. No. 1.3e-07;
Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
DB 527 TCCTGGCTCTGTGTGTCTGTGTCTGTGTCTGTCT 108
|||||TCCTGGCTCTGTGTGTCTGTGTCTGTGTCTGTCT 493

RESULT 7
AO881246/c 529 bp DNA linear GSS 09-NOV-1999
LOCUS HS_5137_B1_F08_T7 RPTI-11 Human Male BAC Library Homo sapiens
DEFINITION genomic clone Plate=8905 Col=15 Row=L, DNA sequence.
ACCESSION AO881246
VERSION AO881246.1 GI:6312713
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 529)
Maitras,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D., and
Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
99380589
Contact: Maitras GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPTI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buflalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buflalo.edu/ordering_bac.htm)
or from Research h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 8905 row: L column: 15
Seq primer: T7
Class: BAC ends
High quality sequence stop: 529.
Location/Qualifiers

FEATURES
Source

1. 529
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=8905 Col=15 Row=L"
/clone_lib="RPTI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from a combination of EcoRI and
and partially digested with a combination of EcoRI and

EcoRI Methylase. Size selected DNA was cloned into the
pBAC3.6 vector at EcoRI sites"
BASE COUNT 137 a 147 c 121 g 117 t 7 others
ORIGIN

Query Match 30.6%; Score 34; DB 17; Length 529;
Best Local Similarity 100.0%; Pred. No. 4.2e-07;
Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 60 GGACTATGTAACCTCTGGCTCTGTGTCTGC 93
|||||GGACTATGTAACCTCTGGCTCTGTGTCTGC 349
Db 382 GGACTATGTAACCTCTGGCTCTGTGTCTGC 349

RESULT 8
AO320567/c 482 bp DNA linear GSS 04-MAY-1999
LOCUS RPTI11-99N1.TV RPTI-11 Homo sapiens genomic clone RPTI-11-99N1, DNA
DEFINITION sequence.
ACCESSION AO320567
VERSION AO320567.1 GI:4050696
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 482)
Adams,M.D., Rounsley,S.D., Zhao,S., Bass,S., Linher,K., Golden,K.,
Berry,K., Granger,D., Suh,E., Wible,C., de Jong,P. and Venter,J.C.
Use of human BAC End Sequences for Sequence-Ready Map Building
Unpublished (1998)
Contact: Shaying Zhao, William Nierman, Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850
Tel: 301 838 0200
Fax: 301 838 0208
Email: hbeet@tigr.org
Clones are derived from the human BAC library RPTI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buflalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buflalo.edu/ordering)
Research Genetics (info@resgen.com). BAC end search page:
http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html
Seq primer: T7
Class: BAC ends.
Location/Qualifiers

FEATURES
Source

1. 482
/organism="Homo sapiens"
/db_xref="GDB:7537944"
/db_xref="taxon:9606"
/clone="RPTI-11-99N1"
/clone_lib="RPTI-11"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPTI11 Human Male BAC Library"
BASE COUNT 142 a 121 c 106 g 112 t 1 others
ORIGIN

Query Match 29.7%; Score 33; DB 17; Length 482;
Best Local Similarity 100.0%; Pred. No. 1.4e-06;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 53 TGGGGCCGGAGTATGTAACCTCTGTGTCT 85
|||||TGGGGCCGGAGTATGTAACCTCTGTGTCT 306
Db 338 TGGGGCCGGAGTATGTAACCTCTGTGTCT 306

RESULT 9
AO420187/c 563 bp DNA linear GSS 23-MAR-1999
LOCUS RPTI-11-185J19.TV RPTI-11 Homo sapiens genomic clone RPTI-11-185J19
DEFINITION

, DNA sequence.
 ACCESSION AQ420187
 VERSION AQ420187.1 GI:4477911
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 563)
 AUTHORS Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter ,J.C.
 TITLE Use of BAC end Sequences from Library RPCI-11 for Sequence-Ready Map Building
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI-11-185J19.TV
 Contact: Shaying Zhao, William Niernan, Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbeetlgr.org
 Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html.
 Seq primer: SP6
 Class: BAC ends.
 FEATURES
 source
 Location/Qualifiers
 1..563
 /organism="Homo sapiens"
 /db_xref="GDB:7570890"
 /db_xref="taxon:9606"
 /clone="RPCI-11-185J19"
 /clone_1lb="RPCI-11"
 /sex="Male"
 /cell_type="Lymphocytes"
 /note="Vector: pBACe3.6; Site.1: EcoRI; Site.2: EcoRI; RPCI11 Human Male BAC Library"
 BASE COUNT 173 a 127 c 114 g 149 t
 ORIGIN
 Query Match 29.7%; Score 33; DB 17; Length 563;
 Best Local Similarity 100.0%; Pred. No. 1.4e-06;
 Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 53 TGGGGCCGAGTATGTAAACCTCGGTCCTCT 85
 ||||||||||||||||||||||||||||||||
 333 TGGGGCCGAGTATGTAAACCTCGGTCCTCT 301
 RESULT 10
 AQ386439/c 723 bp DNA linear GSS 21-MAY-1999
 LOCUS RPCI11-154D6.TJ RPCI-11 Homo sapiens genomic clone RPCI-11-154D6,
 DEFINITION DNA sequence.
 ACCESSION AQ386439
 VERSION AQ386439.1 GI:4357462
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 723)
 AUTHORS Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter ,J.C.
 TITLE Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready Map Building
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI11-154D6.TV
 Contact: Shaying Zhao, William Niernan, Mark Adams

Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbeetlgr.org
 Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html.
 Seq primer: SP6
 Class: BAC ends.
 FEATURES
 source
 Location/Qualifiers
 1..723
 /organism="Homo sapiens"
 /db_xref="GDB:7558829"
 /db_xref="taxon:9606"
 /clone="RPCI-11-154D6"
 /clone_1lb="RPCI-11"
 /sex="Male"
 /cell_type="Lymphocytes"
 /note="Vector: pBACe3.6; Site.1: EcoRI; Site.2: EcoRI; RPCI11 Human Male BAC Library"
 BASE COUNT 224 a 165 c 135 g 199 t
 ORIGIN
 Query Match 29.7%; Score 33; DB 17; Length 723;
 Best Local Similarity 100.0%; Pred. No. 1.5e-06;
 Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 53 TGGGGCCGAGTATGTAAACCTCGGTCCTCT 85
 ||||||||||||||||||||||||||||||||
 Db 331 TGGGGCCGAGTATGTAAACCTCGGTCCTCT 299
 RESULT 11
 A2521751 553 bp DNA linear GSS 16-OCT-2000
 LOCUS RPCI-11-175G22.TVB RPCI-11 Homo sapiens genomic clone
 DEFINITION RPCI-11-175G22, DNA sequence.
 ACCESSION A2521751
 VERSION A2521751.1 GI:10834261
 KEYWORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 553)
 AUTHORS Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter ,J.C.
 TITLE BAC end sequences of library RPCI-11
 JOURNAL Unpublished (1997)
 COMMENT Other GSSs: RPCI-11-175G22.TJ
 Contact: Shaying Zhao
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850, USA
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: szhao@tigr.org
 Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genet cs (info@resgen.com). BAC end search page: http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html.
 This BAC end was generated during the R&D process and may have higher chance of clone tracking errors.
 Seq primer: T7
 Class: BAC ends.
 Location/Qualifiers

```

source
1. .553
/db_xref="Homo sapiens"
/db_xref="GDB:7566981"
/db_xref="taxon:9606"
/clone="RPCI-11-175G22"
/clone_lib="RPCI-11"
/sex="Male"
/cell_type="Lymphocytes"
/notes="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC11 Human Male BAC Library"
BASE COUNT      109 a      148 c      137 g      138 t      1 others
ORIGIN
Query Match      27.9%; Score 31; DB 17; Length 553;
Best Local Similarity 100.0%; Pred. No. 1,6e-05;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 68 TAAACTCCTGGCTCTGTGTGCTGAG 98
140 TAAACTCCTGGCTCTGTGTGCTGAG 170

RESULT 12
AG179297      695 bp      DNA      linear      GSS 09-JAN-2002
LOCUS      Pan troglodytes DNA, clone: RP43-051111.TJ, genomic survey
DEFINITION
ACCESSION      AG179297
VERSION      AG179297
KEYWORDS      GSS.
SOURCE      Pan troglodytes male lymphocytes DNA, clone_lib:RPCI-43 Chimpanzee
      Male BAC library clone:RP43-051111.TJ.
      Pan troglodytes
      Pan troglodytes
ORGANISM      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
      Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Pan.
REFERENCE
AUTHORS      Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
      Totoki,Y., Watanabe,H. and Sakaki,Y.
      BAC end sequences of Library RPCI-43
      Unpublished
      2 (bases 1 to 695)
      Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,
      Totoki,Y., Watanabe,H. and Sakaki,Y.
      Direct Submission
      Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical
      and Chemical Research (RIKEN), Genomic Sciences Center (GSC):
      1-7-22 Suehiro-chou,Tsukumi-ku, Yokohama, Kanagawa 230-0045, Japan
      (E-mail:chimpesgsc.riken.go.jp, URL:http://hgp.9sc.riken.go.jp/,
      Tel:81-45-503-9111, Fax:81-45-503-9170)
      Clones are derived from the chimpanzee BAC library RPCI-43 This BAC
      end was generated during the R&D process and may have higher chance
      of clone tracking errors.
      PRIMERS
      Sequencing: TJ
      LIBRARY
      Vector : pBAC3.6
      R.Site 1 : EcoRI
      R.Site 2 : EcoRI.
      Location/Qualifiers
      1. 695
      /organism="Pan troglodytes"
      /db_xref="taxon:9598"
      /clone="RP43-051111.TJ"
      /sex="male"
      /cell_type="Lymphocytes"
      /clone_lib="RPCI-43 Chimpanzee Male BAC Library"
BASE COUNT      243 a      135 c      122 g      192 t      3 others
ORIGIN
Query Match      27.9%; Score 31; DB 17; Length 695;
Best Local Similarity 100.0%; Pred. No. 1,6e-05;
Matches 31; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

QY 61 GAGTATGTAACCTCGGCTCTGTCTGT 91
171 GAGTATGTAACCTCGGCTCTGTCTGT 201

RESULT 13
AO207172      360 bp      DNA      linear      GSS 18-SEP-1998
LOCUS      HS-3239_B1.C03.T7 CIT Approved Human Genomic Sperm Library D Homo
DEFINITION      sapiens genomic clone Plate=3239 Col=5 Row=F, DNA sequence.
ACCESSION      AO207172
VERSION      AO207172.1 GI:3618377
KEYWORDS      GSS.
SOURCE      human.
ORGANISM      Homo sapiens
      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
      Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
      1 (bases 1 to 360)
      Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
      Kellier,A., Shaker,R., Furlong,J., Young,D., Zhao,S., Adams,M.D. and
      Hood,L.
      Sequence-tagged connectors: A sequence approach to mapping and
      scanning the human genome
      Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
      99380589
      Contact: Mahairas GG, Wallace JC, Hood L
      High Throughput Sequencing Center
      University of Washington
      401 Queen Anne Avenue North, Seattle, WA 98109, USA
      Tel: (206) 616-3618
      Fax: (206) 616-3887
      Email: jwallace@u.washington.edu
      Sequence Tagged Connector
      Plate: 3239 row: F column: 5
      Class: BAC ends
      High quality sequence stop: 360.
      Location/Qualifiers
      1. 360
      /organism="Homo sapiens"
      /db_xref="taxon:9606"
      /clone="Plate=3239 Col=5 Row=F"
      /clone_lib="CIT Approved Human Genomic Sperm Library D"
      /sex="male"
      /note="Organ: sperm; Vector: pBelOAPC11; BAC Clones in
      E-Coli DH10B"
BASE COUNT      63 a      93 c      107 g      93 t      4 others
ORIGIN
Query Match      27.0%; Score 30; DB 17; Length 360;
Best Local Similarity 100.0%; Pred. No. 4.9e-05;
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 79 GGTCTGTGTGCTGCTGAGTGGCTGCT 108
171 GGTCTGTGTGCTGCTGAGTGGCTGCT 200

RESULT 14
AO115544/c      399 bp      DNA      linear      GSS 20-APR-1999
LOCUS      RPC11-57K21.7K RPCI-11 Homo sapiens genomic clone RPCI-11-57K21,
DEFINITION      DNA sequence.
ACCESSION      AO115544
VERSION      AO115544.1 GI:3491665
KEYWORDS      GSS.
SOURCE      human.
ORGANISM      Homo sapiens
      Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
      Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
      1 (bases 1 to 399)
      Adams,M.D., Rounsley,S.D., Zhao,S., Bass,S., Linher,K., Golden,K.,
      Berry,K., Granger,D., Suh,E., Wible,C., de Jong,P. and Venter,J.C.
      Use of human BAC End Sequences for Sequence-Ready Map Building
      TITLE

```

JOURNAL
COMMENT

Unpublished (1998)
Other_GSSs: RPC11-57K21.TJ
Contact: Mark Adams
Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208

Email: mdadams@tigr.org
Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/tldb/hungen/bac_end_search/bac_end_search.html
Class: BAC ends.

FEATURES

source

Location/Qualifiers
1..399
/organism="Homo sapiens"
/db_xref="GDB:7521764"
/db_xref="taxon:9606"
/clone="RPC1-11-57K21"
/clone_11b="RPC1-11"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC11 Human Male BAC Library"
BASE COUNT 120 a 111 c 95 g 73 t
ORIGIN

Query Match

Best Local Similarity 100.0%; Score 30; DB 17; Length 399;
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCGGTCTCTGTGTG 90
|||||
Db 380 GAGTATGTAACCTCGGTCTCTGTGTG 351

RESULT 15
A0116061/c

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

COMMENT

/organism="Homo sapiens"
/db_xref="GDB:7521786"
/db_xref="taxon:9606"
/clone="RPC1-11-57L19"
/clone_11b="RPC1-11"
/sex="Male"
/cell_type="Lymphocytes"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC11 Human Male BAC Library"
BASE COUNT 131 a 115 c 103 g 86 t
ORIGIN

BASE COUNT

131 a 115 c 103 g 86 t

ORIGIN

Query Match

Best Local Similarity 100.0%; Score 30; DB 17; Length 435;
Matches 30; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCGGTCTCTGTGTG 90
|||||
Db 379 GAGTATGTAACCTCGGTCTCTGTGTG 350

Search completed: April 25, 2003, 00:52:48
Job time : 378.284 secs


```

: MOLECULE TYPE: DNA (genomic)
: HYPOTHETICAL: NO
: ANTI-SENSE: UNKNOWN
: ORIGINAL SOURCE:
: ORGANISM: PORPHYROMONAS GINGIVALIS
: FEATURE:
: NAME/KEY: misc-feature
: LOCATION: 1...536
: US-09-221-017B-1081

Query Match      14.4% Score 16: DB 4; Length 536;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 26 ATCTTTTCATCTT 41
    |||||
Db 287 ATCTTTTCATCTT 272

>
> 09-149-476-307
> Sequence 307, Application US/09149476
> Patent No. 6420526
> GENERAL INFORMATION:
> APPLICANT: Rosen et al.
> TITLE OF INVENTION: 186 Human Secreted proteins
> FILE REFERENCE: P2002P1
> CURRENT APPLICATION NUMBER: US/09/149,476
> EARLIER APPLICATION NUMBER: 1998-09-08
> EARLIER APPLICATION NUMBER: PCT/US98/04493
> EARLIER FILING DATE: 1998-03-06
> EARLIER APPLICATION NUMBER: 60/040,162
> EARLIER FILING DATE: 1997-03-07
> EARLIER APPLICATION NUMBER: 60/040,333
> EARLIER FILING DATE: 1997-03-07
> EARLIER APPLICATION NUMBER: 60/038,621
> EARLIER FILING DATE: 1997-03-07
> EARLIER APPLICATION NUMBER: 60/040,626
> EARLIER FILING DATE: 1997-03-07
> EARLIER APPLICATION NUMBER: 60/040,334
> EARLIER FILING DATE: 1997-03-07
> EARLIER APPLICATION NUMBER: 60/040,336
> EARLIER FILING DATE: 1997-03-07
> EARLIER APPLICATION NUMBER: 60/040,163
> EARLIER FILING DATE: 1997-03-07
> EARLIER APPLICATION NUMBER: 60/047,600
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,615
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,597
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,502
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,633
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,583
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,617
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,618
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,503
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,592
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,581
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,584
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,500
> EARLIER FILING DATE: 1997-05-23
> EARLIER APPLICATION NUMBER: 60/047,587
> EARLIER FILING DATE: 1997-05-23

: EARLIER APPLICATION NUMBER: 60/047,492
: EARLIER FILING DATE: 1997-05-23
: EARLIER APPLICATION NUMBER: 60/047,598
: EARLIER FILING DATE: 1997-05-23
: EARLIER APPLICATION NUMBER: 60/047,613
: EARLIER FILING DATE: 1997-05-23
: EARLIER APPLICATION NUMBER: 60/047,582
: EARLIER FILING DATE: 1997-05-23
: EARLIER APPLICATION NUMBER: 60/047,596
: EARLIER FILING DATE: 1997-05-23
: EARLIER APPLICATION NUMBER: 60/047,612
: EARLIER FILING DATE: 1997-05-23
: EARLIER APPLICATION NUMBER: 60/047,632
: EARLIER FILING DATE: 1997-05-23
: EARLIER APPLICATION NUMBER: 60/047,601
: EARLIER FILING DATE: 1997-05-23
: EARLIER APPLICATION NUMBER: 60/043,580
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,568
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,314
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,569
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,311
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,671
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,674
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,669
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,312
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,313
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,672
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/043,315
: EARLIER FILING DATE: 1997-04-11
: EARLIER APPLICATION NUMBER: 60/048,974
: EARLIER FILING DATE: 1997-06-06
: EARLIER APPLICATION NUMBER: 60/056,886
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,877
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,889
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,893
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,630
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,878
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,662
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,872
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,882
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,637
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,903
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,888
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,879
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,880
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,894
: EARLIER FILING DATE: 1997-08-22
: EARLIER APPLICATION NUMBER: 60/056,911

```

```

; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,636
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,874
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,910
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,864
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,631
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,845
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,892
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/057,761
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/047,595
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,599
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,588
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,585
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,586
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,590
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,594
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,589
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,593
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/047,614
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/043,578
; EARLIER FILING DATE: 1997-04-11
; EARLIER APPLICATION NUMBER: 60/043,576
; EARLIER FILING DATE: 1997-04-11
; EARLIER APPLICATION NUMBER: 60/047,501
; EARLIER FILING DATE: 1997-05-23
; EARLIER APPLICATION NUMBER: 60/043,670
; EARLIER FILING DATE: 1997-04-11
; EARLIER APPLICATION NUMBER: 60/056,632
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,664
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,876
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,881
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,909
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,875
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,862
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,887
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/056,908
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/048,964
; EARLIER FILING DATE: 1997-06-06
; EARLIER APPLICATION NUMBER: 60/057,650
; EARLIER FILING DATE: 1997-09-05
; EARLIER APPLICATION NUMBER: 60/056,884
; EARLIER FILING DATE: 1997-08-22
; EARLIER APPLICATION NUMBER: 60/057,669
; EARLIER FILING DATE: 1997-09-05
; EARLIER APPLICATION NUMBER: 60/049,610
; EARLIER FILING DATE: 1997-06-13

```

```

; EARLIER APPLICATION NUMBER: 60/061,060
; EARLIER FILING DATE: 1997-10-02

```

```

Query Match 14.4%; Score 16; DB 4; Length 997;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Oy 8 GATCTTTGCTTGA 23
|||||
Db 662 GATCTTTGCTTGA 677

```

```

RESULT 3
US-09-233-506-1
; Sequence 1, Application US/09233506
; Patent No. 6136580

```

```

; GENERAL INFORMATION:
; APPLICANT: Fukuda, Minoru

```

```

; TITLE OF INVENTION: A Beta-1-6-N-Acetylglucosaminyltransferase That Forms

```

```

; FILE REFERENCE: P-1J 3415
; CURRENT APPLICATION NUMBER: US/09/233,506

```

```

; CURRENT FILING DATE: 1999-01-19
; NUMBER OF SEQ ID NOS: 14

```

```

; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1

```

```

; LENGTH: 2128
; TYPE: DNA

```

```

; ORGANISM: Homo sapiens
; FEATURE:

```

```

; NAME/KEY: CDS
; LOCATION: (354)..(1670)

```

```

US-09-233-506-1

```

```

Query Match 14.4%; Score 16; DB 3; Length 2128;
Best Local Similarity 100.0%; Pred. No. 13;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Oy 8 GATCTTTGCTTGA 23
|||||
Db 1821 GATCTTTGCTTGA 1836

```

```

RESULT 4
US-09-334-601-6/c
; Sequence 6, Application US/09334601
; Patent No. 6280989

```

```

; GENERAL INFORMATION:
; APPLICANT: Kapitonov, Dmitri

```

```

; TITLE OF INVENTION: NOVEL SLALYTRANSFERASES
; FILE REFERENCE: VCUIP-6

```

```

; CURRENT APPLICATION NUMBER: US/09/334,601
; CURRENT FILING DATE: 1999-06-17

```

```

; NUMBER OF SEQ ID NOS: 94
; SOFTWARE: PatentIn Ver. 2.0

```

```

; SEQ ID NO 6
; LENGTH: 2178

```

```

; TYPE: DNA
; ORGANISM: Homo sapiens

```

```

US-09-334-601-6

```

```

Query Match 14.4%; Score 16; DB 4; Length 2178;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

Oy 26 ATCTTTTTCATCTTT 41
|||||
Db 2150 ATCTTTTTCATCTTT 2135

```

```

RESULT 5

```

US-09-334-601-1/C
; Sequence 1, Application US/09334601
; Patent No. 6280989
; GENERAL INFORMATION:
; APPLICANT: Kapitonov, Dmitri
; APPLICANT: Yu, Robert
; TITLE OF INVENTION: NOVEL STALYLTRANSFERASES
; FILE REFERENCE: VCUJP-6
; CURRENT APPLICATION NUMBER: US/09/334,601
; CURRENT FILING DATE: 1999-06-17
; NUMBER OF SEQ ID NOS: 94
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1
; LENGTH: 2288
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (29)..(1282)
; 09-334-601-1

Query Match 14.4%; Score 16; DB 4; Length 2288;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 26 ATCTTTTCATCTT 41
|||||
DB 2260 ATCTTTTCATCTT 2245

RESULT 6
US-07-746-705A-16
; Sequence 16, Application US/07746705A
; Patent No. 5451516
; GENERAL INFORMATION:
; APPLICANT: Matthews, Benjamin F.
; APPLICANT: Weismann, Jane M.
; TITLE OF INVENTION: A Recombinant DNA Molecule Encoding
; TITLE OF INVENTION: a Bifunctional Plant Enzyme: Aspartokinase and Homoserine
; NUMBER OF SEQUENCES: 16
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle S. Graeter
; STREET: Bldg. 005, Room 402, BARC-W
; CITY: Beltsville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20705
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/07/746,705A
; FILING DATE: 19910816
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Graeter, Janelle S.
; REGISTRATION NUMBER: 35,024
; REFERENCE/DOCKET NUMBER: 4000.91
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (301)504-5676
; TELEFAX: (301)504-5060
; INFORMATION FOR SEQ ID NO: 16:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2915 base pairs
; TYPE: NUCLEIC ACID
; STRANDEDNESS: both
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA to mRNA
; HYPOTHETICAL: NO
; ANTI-SENSE: NO

FEATURE:
; NAME/KEY: CDS
; LOCATION: 2..2593
; 09-334-601-1

Query Match 14.4%; Score 16; DB 1; Length 2915;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 38 CTTTGACGAGCTTCT 53
|||||
DB 1977 CTTTGACGAGCTTCT 1992

RESULT 7
US-08-380-182-18
; Sequence 18, Application US/08380182
; Patent No. 5858749
; GENERAL INFORMATION:
; APPLICANT: Matthews, Benjamin F.
; APPLICANT: Weismann, Jane M.
; TITLE OF INVENTION: A Bifunctional Protein From Carrots
; TITLE OF INVENTION: (Daucus carota) with Aspartokinase and Homoserine
; NUMBER OF SEQUENCES: 23
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Janelle S. Graeter
; STREET: Room 411, Bldg. 005, BARC-W
; CITY: Beltsville
; STATE: Maryland
; COUNTRY: USA
; ZIP: 20705
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/380,182
; FILING DATE:
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Graeter, Janelle S.
; REGISTRATION NUMBER: 35,024
; REFERENCE/DOCKET NUMBER: 0226.94
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 301-504-6629
; TELEFAX: 301-504-5060
; INFORMATION FOR SEQ ID NO: 18:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2915 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Daucus carota
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 2..2593
; 08-380-182-18

Query Match 14.4%; Score 16; DB 2; Length 2915;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 38 CTTTGACGAGCTTCT 53
|||||
DB 1977 CTTTGACGAGCTTCT 1992

RESULT 8
US-09-334-601-5/c
Sequence 5, Application US/09334601
Patent No. 6280989
GENERAL INFORMATION:
APPLICANT: Kapitonov, Dmitri
APPLICANT: Yu, Robert
TITLE OF INVENTION: NOVEL STALYLTRANSFERASES
FILE REFERENCE: YCUIP-6
CURRENT APPLICATION NUMBER: US/09/334,601
CURRENT FILING DATE: 1999-06-17
NUMBER OF SEQ ID NOS: 94
SOFTWARE: Patentln Ver. 2.0
SEQ ID NO 5
LENGTH: 3494
TYPE: DNA
ORGANISM: Homo sapiens
US-09-334-601-5

Query Match 14.4%; Score 16; DB 4; Length 3494;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 26 ATCTTTTCATCTT 41
|||||
DB 3467 ATCTTTTCATCTT 3452

RESULT 9
US-09-024-020B-1/c
Sequence 1, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLYVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-3322
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 5977 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear

MOLECULE TYPE: DNA (genomic)
US-09-024-020B-1

Query Match 14.4%; Score 16; DB 3; Length 5977;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 80 GTCTGTGTGTGCTT 95
|||||
DB 5814 GTCTGTGTGTGCTT 5799

RESULT 10
US-09-425-043-1/c
Sequence 1, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLYVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentln Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-3322
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 5977 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-425-043-1

Query Match 14.4%; Score 16; DB 4; Length 5977;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 80 GTCTGTGTGTGCTT 95
|||||
DB 5814 GTCTGTGTGTGCTT 5799

RESULT 11
US-09-024-020B-2/c

Sequence 2, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESSES:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: FLOPPY disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 6007 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-024-020B-2

Query Match 14.4%; Score 16; DB 3; Length 6007;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

DB 5844 GTCTCTGTGTGCTCT 5829

QY 80 GTCTCTGTGTGCTCT 95
|||||

RESULT 12
US-09-425-043-2/c
Sequence 2, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESSES:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA

COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: FLOPPY disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 09/024,020
FILING DATE: 16-FEB-1998
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 6007 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-425-043-2

Query Match 14.4%; Score 16; DB 4; Length 6007;
Best Local Similarity 100.0%; Pred. No. 15;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 80 GTCTCTGTGTGCTCT 95
|||||

DB 5844 GTCTCTGTGTGCTCT 5829

RESULT 13
US-09-024-020B-7/c
Sequence 7, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESSES:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: FLOPPY disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997

```

ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 6556 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-024-020B-7

Query Match
Best Local Similarity 14.4%; Score 16; DB 3; Length 6556;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

80 GTCTCTGTGTGTCCT 95
DB 5961 GTCTCTGTGTGTCCT 5946

RESULT 14
US-09-425-043-7/c
Sequence 7, Application US/09425043
Patent No. 6335172
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/425,043
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 6556 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)

```

```

US-09-425-043-7
Query Match
Best Local Similarity 14.4%; Score 16; DB 4; Length 6556;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

80 GTCTCTGTGTGTCCT 95
DB 5961 GTCTCTGTGTGTCCT 5946

RESULT 15
US-09-024-020B-43/c
Sequence 43, Application US/09024020B
Patent No. 6030810
GENERAL INFORMATION:
APPLICANT: DELGADO, STEPHEN G.
APPLICANT: DIETRICH, PAUL S.
APPLICANT: FISH, LINDA M.
APPLICANT: HERMAN, RONALD C.
APPLICANT: SANGAMESWARAN, LAKSHMI
TITLE OF INVENTION: NOVEL CLONED TETRODOTOXIN-SENSITIVE
TITLE OF INVENTION: SODIUM CHANNEL I-SUBUNIT AND A SPLICE VARIANT THEREOF
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: JANET PAULINE CLARK
STREET: 3401 HILLVIEW AVENUE, MS A2-250
CITY: PALO ALTO
STATE: CA
COUNTRY: U.S.A.
ZIP: 94304-1397
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/024,020B
FILING DATE: 16-FEB-1998
CLASSIFICATION: 536
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/039,447
FILING DATE: 26-FEB-1997
ATTORNEY/AGENT INFORMATION:
NAME: CLARK, JANET P.
REGISTRATION NUMBER: 34,799
REFERENCE/DOCKET NUMBER: R0020B-REG
TELECOMMUNICATION INFORMATION:
TELEPHONE: (650) 852-3097
TELEFAX: (650) 855-5322
INFORMATION FOR SEQ ID NO: 43:
SEQUENCE CHARACTERISTICS:
LENGTH: 6586 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-09-024-020B-43

Query Match
Best Local Similarity 14.4%; Score 16; DB 3; Length 6586;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

80 GTCTCTGTGTGTCCT 95
DB 5991 GTCTCTGTGTGTCCT 5976

```

Search completed: April 25, 2003, 00:54:06
Job time : 19.1361 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 00:41:49 ; Search time 22.7692 Seconds

(without alignments)
5304.628 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggtgagcttcttgcctt.....gcctgagtgctctact 111

Scoring table: OLIGO_NUC
Gapop 60.0, Gapext 60.0

Searched: 709820 seqs, 544064369 residues

Database size: 0
Total number of hits satisfying chosen parameters: 1419640

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database:

Published Applications_NA:*

- 1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq:*
- 2: /cgn2_6/ptodata/1/pubpna/PCOT_NEW_PUB.seq:*
- 3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq:*
- 4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq:*
- 5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq:*
- 6: /cgn2_6/ptodata/1/pubpna/PCOTUS_PUBCOMB.seq:*
- 7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq:*
- 8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq:*
- 9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
- 10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq:*
- 11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*
- 12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq:*
- 13: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
- 14: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	33	29.7	684973	10 US-09-263-959-1	Sequence 1, Appli
2	18	16.2	431	10 US-09-866-562-80	Sequence 80, Appl
3	18	16.2	431	10 US-09-866-562-87	Sequence 87, Appl
4	18	16.2	570	10 US-09-864-761-9118	Sequence 9118, Ap
5	17	15.3	531	9 US-10-092-154-1899	Sequence 1899, Ap
6	17	15.3	531	10 US-09-764-847-1899	Sequence 1899, Ap
7	17	15.3	609	10 US-09-974-300-8084	Sequence 8084, Ap
8	17	15.3	1047	10 US-09-822-830A-468	Sequence 468, Ap
9	17	15.3	172637	10 US-09-805-458A-3	Sequence 458, Ap
10	17	15.3	1503841	9 US-09-946-807-1	Sequence 3, Appli
11	17	15.3	1503841	10 US-09-795-668-1	Sequence 1, Appli
12	17	15.3	1503841	10 US-09-795-668-1	Sequence 1, Appli
13	16	14.4	281	10 US-09-964-824A-385	Sequence 385, App
14	16	14.4	374	9 US-10-046-935-2033	Sequence 2033, Ap
15	16	14.4	374	9 US-09-878-178-2033	Sequence 2033, Ap
16	16	14.4	374	9 US-10-146-502-2033	Sequence 2033, Ap
17	16	14.4	374	9 US-10-060-036-985	Sequence 985, App
18	16	14.4	374	9 US-10-060-036-2528	Sequence 2528, Ap
19	16	14.4	436	9 US-09-918-995-26108	Sequence 26108, A

C 20	16	14.4	447	10 US-09-880-107-871	Sequence 871, App
C 21	16	14.4	481	10 US-09-560-863-45	Sequence 45, Appl
C 22	16	14.4	494	9 US-09-918-995-20796	Sequence 20796, A
C 23	16	14.4	497	10 US-09-864-761-1356	Sequence 1356, Ap
C 24	16	14.4	524	10 US-09-797-207-5	Sequence 5, Appli
C 25	16	14.4	548	10 US-09-864-761-12421	Sequence 12421, A
C 26	16	14.4	552	10 US-09-998-598-1325	Sequence 1325, Ap
C 27	16	14.4	557	10 US-09-764-877-746	Sequence 746, App
C 28	16	14.4	557	10 US-09-764-877-3495	Sequence 3495, Ap
C 29	16	14.4	747	12 US-10-001-879-101	Sequence 101, App
C 30	16	14.4	805	9 US-09-984-245-106	Sequence 106, App
C 31	16	14.4	805	9 US-09-966-262-106	Sequence 106, App
C 32	16	14.4	805	9 US-09-983-966-106	Sequence 106, App
C 33	16	14.4	805	9 US-10-143-090-106	Sequence 106, App
C 34	16	14.4	997	9 US-09-809-391-307	Sequence 307, App
C 35	16	14.4	1009	10 US-09-764-864-12	Sequence 12, Appl
C 36	16	14.4	2002	10 US-09-925-300-592	Sequence 592, App
C 37	16	14.4	2017	9 US-10-102-806-137	Sequence 137, App
C 38	16	14.4	2108	10 US-09-797-207-3	Sequence 3, Appli
C 39	16	14.4	2147	9 US-09-981-353-43	Sequence 43, Appl
C 40	16	14.4	2229	10 US-09-925-297-337	Sequence 337, App
C 41	16	14.4	2268	9 US-10-102-806-142	Sequence 142, App
C 42	16	14.4	2287	10 US-09-764-864-473	Sequence 473, App
C 43	16	14.4	2319	10 US-09-874-390-1	Sequence 1, Appli
C 44	16	14.4	2340	9 US-09-964-245-51	Sequence 51, Appl
C 45	16	14.4	2340	9 US-09-966-262-51	Sequence 51, Appl

ALIGNMENTS

RESULT 1
US-09-263-959-1/c
Sequence 1, Application US/09263959
Patent No. US20020150891A1
GENERAL INFORMATION:
APPLICANT: Hood, Leroy E.
APPLICANT: Rowen, Lee
TITLE OF INVENTION: DIAGNOSTIC AND THERAPEUTIC COMPOSITIONS AND METHODS WHICH U
NUMBER OF SEQUENCES: 1279
CORRESPONDENCE ADDRESS:
ADDRESSEE: Seed and Berry LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: US
ZIP: 98104-7092
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/263,959
FILING DATE: 05-MAR-1999
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Mcmasters, David D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 920010.426C2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 684973 base pairs
TYPE: nucleic acid
STRANDEDNESS: Single
TOPOLOGY: linear
US-09-263-959-1
Query Match 29.7%; Score 33; DB 10; Length 684973;

Best Local Similarity 100.0%; Pred. No. 2.4e-08;
Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 66 TGTAAACTCCTGGGTCCTGTGTGTCCTGAG 98
|||||
Db 404690 TGTAAACTCCTGGGTCCTGTGTGTCCTGAG 404658

RESULT 2
US-09-866-562-80

Sequence 80, Application us/09866562
Patent No. US20020009758a1
GENERAL INFORMATION:
APPLICANT: Harlocker, Susan L.
APPLICANT: Wang, Tongtong
APPLICANT: Bangur, Chaitanya S.
APPLICANT: Klee, Jennifer
APPLICANT: Switzer, Anne
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
OF LUNG CANCER.
TITLE OF INVENTION: AND DIAGNOSIS OF LUNG CANCER.
FILE REFERENCE: 210121_502
CURRENT APPLICATION NUMBER: US/09/866,562
CURRENT FILING DATE: 2001-05-25
NUMBER OF SEQ ID NOS: 96

```

: ORGANISM: Homo sapiens
:
: FEATURE:
:
: NAME/KEY: misc_feature
: LOCATION: 361..431
:
: OTHER INFORMATION: n = A,T,C or G
US-09-866-562-80

```

Query Match	16.28;	Score 18;	DB 10;	Length 431;
Best [local] similarity	100.08;	Pred NC	2;	

Best Local Similarity 100.0%; Pred. NO. 3;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0

oy	91	TGCCTGAGTGGCTGCTCT	108
db	18	TGCCTGAGTGGCTGCTCT	35

RESULT 3
US-09-866-562-87
; Sequence 87, Application US/09866562
; Patent No. US2002009758A1
GENERAL INFORMATION

APPLICANT: Harlocker, Susan L.
APPLICANT: Wang, Tongtong
APPLICANT: Bangut, Chaitanya S.
APPLICANT: Klee, Jennifer
APPLICANT: Switzer, Anne
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
TITLE OF INVENTION: AND DIAGNOSIS OF LUNG CANCER.

```

; FILE REFERENCE: 210121.502
; CURRENT APPLICATION NUMBER: US/09/866,562
; CURRENT FILING DATE: 2001-05-25
; NUMBER OF SEQ ID NOS: 96

```

```

: SEQ ID NO 87
: LENGTH: 431
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: 361..431
: OTHER INFORMATION: n = A,T,C or G
US-09-866-562-87

```

Query Match	16.2%;	Score 18;	DB 10;	Length 431;
Best Local Similarity	100.0%;	Pred. No. 3;		
Matches 18;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0

OY	91	TGCCTGAGTGGCTGCTCT	108
Db	18	TGCCTGAGTGGCTGCTCT	35

RESULT 4
US-09-864-761-9118/c

Patent No. US20020048763A1
GENERAL INFORMATION:

APPLICANT: Penn, Sharon G.
 APPLICANT: Rank, David R.
 APPLICANT: Hanzel, David K.
 APPLICANT: Chen, Wensheng
 TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
 TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY
 FILE REFERENCE: Aecomica-X-1

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65
66
67
68
69
70
71
72
73
74
75
76
77
78
79
80
81
82
83
84
85
86
87
88
89
90
91
92
93
94
95
96
97
98
99
100
101
102
103
104
105
106
107
108
109
110
111
112
113
114
115
116
117
118
119
120
121
122
123
124
125
126
127
128
129
130
131
132
133
134
135
136
137
138
139
140
141
142
143
144
145
146
147
148
149
150
151
152
153
154
155
156
157
158
159
160
161
162
163
164
165
166
167
168
169
170
171
172
173
174
175
176
177
178
179
180
181
182
183
184
185
186
187
188
189
190
191
192
193
194
195
196
197
198
199
200
201
202
203
204
205
206
207
208
209
210
211
212
213
214
215
216
217
218
219
220
221
222
223
224
225
226
227
228
229
230
231
232
233
234
235
236
237
238
239
240
241
242
243
244
245
246
247
248
249
250
251
252
253
254
255
256
257
258
259
260
261
262
263
264
265
266
267
268
269
270
271
272
273
274
275
276
277
278
279
280
281
282
283
284
285
286
287
288
289
290
291
292
293
294
295
296
297
298
299
300
301
302
303
304
305
306
307
308
309
310
311
312
313
314
315
316
317
318
319
320
321
322
323
324
325
326
327
328
329
330
331
332
333
334
335
336
337
338
339
340
341
342
343
344
345
346
347
348
349
350
351
352
353
354
355
356
357
358
359
360
361
362
363
364
365
366
367
368
369
370
371
372
373
374
375
376
377
378
379
380
381
382
383
384
385
386
387
388
389
390
391
392
393
394
395
396
397
398
399
400
401
402
403
404
405
406
407
408
409
410
411
412
413
414
415
416
417
418
419
420
421
422
423
424
425
426
427
428
429
430
431
432
433
434
435
436
437
438
439
440
441
442
443
444
445
446
447
448
449
450
451
452
453
454
455
456
457
458
459
460
461
462
463
464
465
466
467
468
469
470
471
472
473
474
475
476
477
478
479
480
481
482
483
484
485
486
487
488
489
490
491
492
493
494
495
496
497
498
499
500
501
502
503
504
505
506
507
508
509
510
511
512
513
514
515
516
517
518
519
520
521
522
523
524
525
526
527
528
529
530
531
532
533
534
535
536
537
538
539
540
541
542
543
544
545
546
547
548
549
550
551
552
553
554
555
556
557
558
559
560
561
562
563
564
565
566
567
568
569
570
571
572
573
574
575
576
577
578
579
580
581
582
583
584
585
586
587
588
589
590
591
592
593
594
595
596
597
598
599
600
601
602
603
604
605
606
607
608
609
610
611
612
613
614
615
616
617
618
619
620
621
622
623
624
625
626
627
628
629
630
631
632
633
634
635
636
637
638
639
640
641
642
643
644
645
646
647
648
649
650
651
652
653
654
655
656
657
658
659
660
661
662
663
664
665
666
667
668
669
670
671
672
673
674
675
676
677
678
679
680
681
682
683
684
685
686
687
688
689
690
691
692
693
694
695
696
697
698
699
700
701
702
703
704
705
706
707
708
709
710
711
712
713
714
715
716
717
718
719
720
721
722
723
724
725
726
727
728
729
730
731
732
733
734
735
736
737
738
739
740
741
742
743
744
745
746
747
748
749
750
751
752
753
754
755
756
757
758
759
760
761
762
763
764
765
766
767
768
769
770
771
772
773
774
775
776
777
778
779
780
781
782
783
784
785
786
787
788
789
790
791
792
793
794
795
796
797
798
799
800
801
802
803
804
805
806
807
808
809
810
811
812
813
814
815
816
817
818
819
820
821
822
823
824
825
826
827
828
829
830
831
832
833
834
835
836
837
838
839
840
84

PRIOR FILING DATE: 2000-10-04
 PRIOR APPLICATION NUMBER: US 60/236,359
 PRIOR FILING DATE: 2000-09-27
 PRIOR APPLICATION NUMBER: PCT/US01/00666
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00667
 PRIOR FILING DATE: 2001-01-30

PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00669
 ; PRIOR FILING DATE: 2001-01-30

PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30

; PRIOR APPLICATION NUMBER: PCT/US01/00663
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00662
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00661
 ; PRIOR FILING DATE: 2001-01-30

PRIOR APPLICATION NUMBER: PCT/US01/006670
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: US 60,234,687
PRIOR FILING DATE: 2000-09-21
PRIOR APPLICATION NUMBER: US 09/608,408
PRIOR FILING DATE: 2000-06-30
PRIOR APPLICATION NUMBER: US 09/774,203
PRIOR FILING DATE: 2001-01-29

```

NUMBER OF SEQ ID NOS: 49117
SOFTWARE: Annomax Sequence Listing Engine vers. 1.1.1
SEQ ID NO 9118
LENGTH: 570

```

```

? TYPE: DNA
? ORGANISM: Homo sapiens
? FEATURE:
? OTHER INFORMATION: MAP TO AF000053.1
? OTHER INFORMATION: EXPRESSED IN HEILA, SIGNAL = 2.8
? OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 2.4
? OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 4.8
? OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL =
? OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 2.3
? OTHER INFORMATION: EXPRESSED IN SPLEEN, SIGNAL = 4

```

OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 3.6	
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 4.4	
US-09-864-761-9118	

Query Match 16.2%; Score 18; DB 10; Length 570;
Best Local Similarity 100.0%; Pred. No. 3;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 85 TGTGTGCTGCTGAGTGGC 102
|||||
Db 505 TGTGTGCTGCTGAGTGGC 488

RESULT 5
US-10-092-154-1899/c
; Sequence 1899, Application US/10092154
; Publication No. US20030054375a1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC009C1
; CURRENT APPLICATION NUMBER: US/10/092,154
; CURRENT FILING DATE: 2002-03-07
; NUMBER OF SEQ ID NOS: 2003
; Prior Application removed - See File Wrapper or Palm
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO: 1899
; LENGTH: 531
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-092-154-1899

Query Match 15.3%; Score 17; DB 9; Length 531;
Best Local Similarity 100.0%; Pred. No. 10;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 28 TCTTTTCATCTTTGCA 44
|||||
Db 141 TCTTTTCATCTTTGCA 125

RESULT 6
US-09-764-847-1899/c
; Sequence 1899, Application US/09764847
; Patent No. US20020132767A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC009
; CURRENT APPLICATION NUMBER: US/09/764,847
; CURRENT FILING DATE: 2001-01-17
; Prior Application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 2003
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO: 1899
; LENGTH: 531
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-847-1899

Query Match 15.3%; Score 17; DB 10; Length 531;
Best Local Similarity 100.0%; Pred. No. 10;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 28 TCTTTTCATCTTTGCA 44
|||||
Db 141 TCTTTTCATCTTTGCA 125

RESULT 7
US-09-974-300-8084/c
; Sequence 8084, Application US/09974300
; Patent No. US20020146721A1
; GENERAL INFORMATION:
; APPLICANT: Berka, Randy M.
; APPLICANT: Clausen, Ib Groth
; TITLE OF INVENTION: Methods For Monitoring Multiple Gene

; TITLE OF INVENTION: Expression
; FILE REFERENCE: 10085,500-US
; CURRENT APPLICATION NUMBER: US/09/974,300
; CURRENT FILING DATE: 2001-10-05
; PRIOR APPLICATION NUMBER: 09/680,598
; PRIOR FILING DATE: 2000-10-06
; PRIOR APPLICATION NUMBER: 60/279,526
; PRIOR FILING DATE: 2001-03-27
; NUMBER OF SEQ ID NOS: 8481
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO: 8084
; LENGTH: 609
; TYPE: DNA
; ORGANISM: Bacillus clausii
US-09-974-300-8084

Query Match 15.3%; Score 17; DB 10; Length 609;
Best Local Similarity 100.0%; Pred. No. 10;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 28 TCTTTTCATCTTTGCA 44
|||||
Db 536 TCTTTTCATCTTTGCA 520

RESULT 8
US-09-822-830A-468
; Sequence 468, Application US/09822830A
; Patent No. US20020142952A1
; GENERAL INFORMATION:
; APPLICANT: Genetics Institute, Inc.
; APPLICANT: Wong, Gordon G.
; APPLICANT: Clark, Hilary
; APPLICANT: Fectel, Kim
; APPLICANT: Agostino, Michael J.
; APPLICANT: Howes, Steven H.
; APPLICANT: Resnick, Richard J.
; APPLICANT: Gulukota, Kamalakara
; APPLICANT: Graham, James R.
; TITLE OF INVENTION: POLYNUCLEOTIDES ENCODING NOVEL SECRETED PROTEINS
; FILE REFERENCE: GIN 6402
; CURRENT APPLICATION NUMBER: US/09/822,830A
; CURRENT FILING DATE: 2001-03-29
; PRIOR APPLICATION NUMBER: 60/195,604
; PRIOR FILING DATE: 2000-04-06
; NUMBER OF SEQ ID NOS: 631
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO: 468
; LENGTH: 1047
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-822-830A-468

Query Match 15.3%; Score 17; DB 10; Length 1047;
Best Local Similarity 100.0%; Pred. No. 10;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 76 CTGGGTCCTGTGTG 92
|||||
Db 828 CTGGGTCCTGTGTG 844

RESULT 9
US-09-805-458A-3
; Sequence 3, Application US/09805458A
; Patent No. US20020042100A1
; GENERAL INFORMATION:
; APPLICANT: YAN, Chunhua et al
; TITLE OF INVENTION: ISOLATED HUMAN ION CHANNEL PROTEINS,
; TITLE OF INVENTION: NUCLEIC ACID MOLECULES ENCODING HUMAN ION CHANNEL PROTEINS,
; TITLE OF INVENTION: AND USBS THEREOF
; FILE REFERENCE: CI000722
; CURRENT APPLICATION NUMBER: US/09/805,458A

```

; CURRENT FILING DATE: 2001-03-14
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 172637
; TYPE: DNA
; ORGANISM: Human
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(172637)
; OTHER INFORMATION: n = A,T,C or G
US-09-805-458A-3
```

```

Query Match          15.3%; Score 17; DB 10; Length 172637;
Best Local Similarity 100.0%; Pred. No. 9,7;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 19 TTGCAGATTCTTTTTC 35
      |||||||
150045 TTGCAGATTCTTTTTC 150061
```

```

RESULT 10
US-09-946-807-1
; Sequence 1, Application US/09946807
; Patent No. US20020165144A1
; GENERAL INFORMATION:
; APPLICANT: Stefansson, Hreinn
; APPLICANT: Steinhorsdottir, Valgerdur
; APPLICANT: Gulcher, Jeffrey R.
; TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
; FILE REFERENCE: 2345.2004-001
; CURRENT APPLICATION NUMBER: US/09/946,807
; CURRENT FILING DATE: 2001-09-05
; PRIOR APPLICATION NUMBER: US/09/795,668
; PRIOR FILING DATE: 2001-02-28
; PRIOR APPLICATION NUMBER: US 09/515,716
; PRIOR FILING DATE: 2000-02-28
; NUMBER OF SEQ ID NOS: 1531
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 1503841
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: y=t/u or c
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: m=a or c
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: k=g or t/u
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: s=g or c
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: w=a or t/u
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: b=g or c or t/u
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: d=a or g or t/u
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
```

```

; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: d=a or g or t/u
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: h=a or c or t/u
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: v=a or g or c
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: n=a or g or c or t/u
US-09-946-807-1
```

```

Query Match          15.3%; Score 17; DB 9; Length 1503841;
Best Local Similarity 100.0%; Pred. No. 9,4;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
QY 26 ATTCTTTTCATCTTTG 42
      |||||||
Db 268726 ATTCTTTTCATCTTTG 268742
```

```

RESULT 11
US-09-795-668-1
; Sequence 1, Application US/09795668
; Patent No. US20020045577A1
; GENERAL INFORMATION:
; APPLICANT: Stefansson, Hreinn
; APPLICANT: Steinhorsdottir, Valgerdur
; APPLICANT: Gulcher, Jeffrey R.
; TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
; FILE REFERENCE: 2345.2004-001
; CURRENT APPLICATION NUMBER: US/09/795,668
; CURRENT FILING DATE: 2001-02-28
; PRIOR APPLICATION NUMBER: US 09/515,716
; PRIOR FILING DATE: 2000-02-28
; NUMBER OF SEQ ID NOS: 1531
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 1503841
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: r=g or a
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: y=t/u or c
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: m=a or c
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: k=g or t/u
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: s=g or c
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: w=a or t/u
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: b=g or c or t/u
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
; OTHER INFORMATION: d=a or g or t/u
; NAME/KEY: misc_feature
; LOCATION: (1)...(1531)
```



```

OTHER INFORMATION: h=a or c or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: v=a or g or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: n=a or g or c or t/u
US-09-795-668-1

```

```

Query Match
Best Local Similarity 15.3%; Score 17; DB 10; Length 1503841;
Pred. No. 9.4;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 26 ATTCTTTTCATCTTTG 42
DB 268726 ATTCTTTTCATCTTTG 268742

```

```

RESULT 12
US-09-795-668-1
Sequence 1, Application US/09795686
Patent No. US20020094954A1
GENERAL INFORMATION:
APPLICANT: Steinhilber, Hrein
APPLICANT: Steinhilber, Valgerdur
APPLICANT: Gulcher, Jeffrey R.
TITLE OF INVENTION: HUMAN SCHIZOPHRENIA GENE
FILE REFERENCE: 2345, 2005-001
CURRENT APPLICATION NUMBER: US/09/795,686
CURRENT FILING DATE: 2001-02-28
PRIOR APPLICATION NUMBER: US 09/515,715
PRIOR FILING DATE: 2000-02-28
NUMBER OF SEQ ID NOS: 1531
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 1
LENGTH: 1503841
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: r=g or a
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: y=l/u or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: m=a or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: k=g or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: s=g or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: w=a or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: b=g or c or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: d=a or g or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: h=a or c or t/u
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: v=a or g or c
NAME/KEY: misc_feature
LOCATION: (1)...(1531)
OTHER INFORMATION: n=a or g or c or t/u
US-09-795-668-1

```

```

Query Match
Best Local Similarity 15.3%; Score 17; DB 10; Length 1503841;
Pred. No. 9.4;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 26 ATTCTTTTCATCTTTG 42
DB 268726 ATTCTTTTCATCTTTG 268742

```

```

RESULT 13
US-09-964-824A-385
Sequence 385, Application US/09964824A
Patent No. US20020102531A1
GENERAL INFORMATION:
APPLICANT: Horigan, Stephen
TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using Sign
FILE REFERENCE: 689290-73
CURRENT APPLICATION NUMBER: US/09/964,824A
CURRENT FILING DATE: 2001-09-27
PRIOR APPLICATION NUMBER: US/60/236,033
PRIOR FILING DATE: 2000-09-28
PRIOR APPLICATION NUMBER: US/60/236,032
PRIOR FILING DATE: 2000-09-28
PRIOR APPLICATION NUMBER: US/60/236,028
PRIOR FILING DATE: 2000-09-28
NUMBER OF SEQ ID NOS: 583
SOFTWARE: Patentin Version 3.0
SEQ ID NO 385
LENGTH: 281
TYPE: DNA
ORGANISM: Homo sapiens
US-09-964-824A-385

```

```

Query Match
Best Local Similarity 14.4%; Score 16; DB 10; Length 281;
Pred. No. 36;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 26 ATTCTTTTCATCTTT 41
DB 22 ATTCTTTTCATCTTT 37

```

```

RESULT 14
US-10-046-935-2033
Sequence 2033, Application US/10046935
Patent No. US20020156011A1
GENERAL INFORMATION:
APPLICANT: Jiang, Yugu
APPLICANT: Harlocker, Susan L.
APPLICANT: Secrist, Heather
APPLICANT: Wang, Aijun
APPLICANT: Stolk, John A.
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
FILE REFERENCE: 210121.527C1
CURRENT APPLICATION NUMBER: US/10/046,935
CURRENT FILING DATE: 2002-01-15
NUMBER OF SEQ ID NOS: 2239
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 2033
LENGTH: 374
TYPE: DNA
ORGANISM: Homo sapiens
US-10-046-935-2033

```

```

Query Match
Best Local Similarity 14.4%; Score 16; DB 9; Length 374;
Pred. No. 36;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

```

```

OY 8 GATCTTTGCCTTGCA 23

```

Db 108 GATCTTTGCCTTGCA 123

RESULT 15

US-09-878-178-2033
 ; Sequence 2033, Application US/09878178
 ; Patent No. US20020177552A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Jiang, Yugu
 ; APPLICANT: Harlocker, Susan L.
 ; APPLICANT: Secrist, Heather
 ; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
 ; TITLE OF INVENTION: AND DIAGNOSIS OF COLON CANCER
 ; FILE REFERENCE: 210121.527
 ; CURRENT APPLICATION NUMBER: US/09/878.178
 ; CURRENT FILING DATE: 2001-06-08
 ; NUMBER OF SEQ ID NOS: 2237
 ; SOFTWARE: fastseq for windows version 4.0
 ; SEQ ID NO 2033
 ; LENGTH: 374
 ; TYPE: DNA
 ; ORGANISM: Homo sapien
 US-09-878-178-2033

Query Match 14.4%; Score 16; DB 9; Length 374;
 Best Local Similarity 100.0%; Pred. No. 36;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Oy 8 GATCTTTGCCTTGCA 23
 |||||
 Db 108 GATCTTTGCCTTGCA 123

Search completed: April 25, 2003, 02:09:14
 Job time : 1043.77 secs

GenCore version 5.1.4.p5_4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:20:48 ; Search time 1096.62 seconds

(without alignments)
10509.347 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_446

Perfect score: 396

Sequence: 1 atgggtgacatttgcctt.....gamctgatatcacttga 396

Scoring table:

Gapop 60.0 , Gapext 60.0

Searched: 2054640 seqs, 14551402878 residues

Size: 0

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database:

GenEmbl:*

1: gb_ba:*

2: gb_hcg:*

3: gb_in:*

4: gb_cm:*

5: gb_ov:*

6: gb_pac:*

7: gb_ph:*

8: gb_pl:*

9: gb_pr:*

10: gb_ro:*

11: gb_sts:*

12: gb_sy:*

13: gb_un:*

14: gb_vl:*

15: em_ba:*

16: em_fun:*

17: em_hum:*

18: em_in:*

19: em_mu:*

20: em_on:*

21: em_or:*

22: em_ov:*

23: em_pat:*

24: em_ph:*

25: em_pl:*

26: em_io:*

27: em_sts:*

28: em_un:*

29: em_vl:*

30: em_hcg_hum:*

31: em_hcg_inv:*

32: em_hcg_other:*

33: em_hcg_mus:*

34: em_hcg_pln:*

35: em_hcg_rod:*

36: em_hcg_mam:*

37: em_hcg_vtl:*

38: em_sy:*

39: em_hcgo_hum:*

40: em_hcgo_mus:*

41: em_hcgo_other:*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	299	75.5	169620	2 AC012674	AC012674 Homo sapi
C 2	48	12.1	143372	9 AL137847	AL137847 Human DNA
C 3	40	10.1	123779	30 AC0221025	AC0221025 Homo sapi
C 4	40	10.1	128118	2 AC076969	AC076969 Homo sapi
C 5	40	10.1	128583	9 AC121249	AC121249 Homo sapi
C 6	40	10.1	148290	9 AC107979	AC107979 Homo sapi
C 7	40	10.1	165649	9 AC103996	AC103996 Homo sapi
C 8	40	10.1	178650	9 AC104303	AC104303 Homo sapi
C 9	40	10.1	192826	9 AC090762	AC090762 Homo sapi
C 10	39	9.8	32918	2 AC007445	AC007445 Homo sapi
C 11	39	9.8	38936	9 AL358817	AL358817 Human DNA
C 12	39	9.8	124271	2 AC025179	AC025179 Homo sapi
C 13	39	9.8	146671	2 AC008814	AC008814 Homo sapi
C 14	39	9.8	159747	2 AP001019	AP001019 Homo sapi
C 15	39	9.8	162740	2 AC034249	AC034249 Homo sapi
C 16	39	9.8	169772	9 AC069538	AC069538 Homo sapi
C 17	39	9.8	175466	9 AL607077	AL607077 Human DNA
C 18	38	9.6	110000	2 AC068875	AC068875 Homo sapi
C 19	38	9.6	166706	9 AC068875	AC068875 Homo sapi
C 20	38	9.6	207408	2 AC068618	AC068618 Homo sapi
C 21	38	9.6	207548	9 AC087283	AC087283 Homo sapi
C 22	37	9.3	135090	9 HS164112	AL009028 Homo sapi
C 23	36	9.1	121720	9 AL581491	AL581491 Human DNA
C 24	35	8.8	153940	9 AC022294	AC022294 Homo sapi
C 25	35	8.8	176253	2 AP001078	AP001078 Homo sapi
C 26	35	8.8	205639	2 AP001793	AP001793 Homo sapi
C 27	35	8.8	210734	2 AC015676	AC015676 Homo sapi
C 28	35	8.8	212055	2 AP000899	AP000899 Homo sapi
C 29	35	8.8	325069	2 AC079737	AC079737 Homo sapi
C 30	34	8.6	99577	9 AC026324	AC026324 Homo sapi
C 31	34	8.6	108040	2 AC068150	AC068150 Homo sapi
C 32	34	8.6	134760	9 AC099484	AC099484 Homo sapi
C 33	34	8.6	146059	2 AC019030	AC019030 Homo sapi
C 34	34	8.6	171347	9 AC099776	AC099776 Homo sapi
C 35	34	8.6	172206	9 AC092119	AC092119 Homo sapi
C 36	34	8.6	172567	2 AC015493	AC015493 Homo sapi
C 37	34	8.6	177447	2 AC104687	AC104687 Homo sapi
C 38	34	8.6	173166	9 AC092375	AC092375 Homo sapi
C 39	34	8.6	182881	9 AC090980	AC090980 Homo sapi
C 40	34	8.6	183016	2 AC068611	AC068611 Homo sapi
C 41	34	8.6	183929	9 AC090797	AC090797 Homo sapi
C 42	34	8.6	273807	2 AC025421	AC025421 Homo sapi
C 43	34	8.6	316296	2 AC092285	AC092285 Homo sapi
C 44	33	8.3	33458	9 HSDJ60101	AL109656 Human DNA
C 45	33	8.3	49616	9 AL365267	AL365267 Human DNA

ALIGNMENTS

RESULT 1

AC012674 169620 bp DNA linear HTG 07-SEP-2000

LOCUS Homo sapiens chromosome 3 clone RP1-458H3, WORKING DRAFT SEQUENCE, 18 unordered pieces.

AC012674

AC012674.10 GI:9719580

VERSION HTG: HTGS PHASE1, HTGS_DRAFT.

KEYWORDS

SOURCE

ORGANISM

Homo sapiens.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

REFERENCE

1 (bases 1 to 169620)

Muzny,D.M., Adams,C., Bailey,M., Barbara,J., Blankenburg,K., Bodots,B., Bouck,J., Bowie,S., Brooks,A., Bunay,C., Bunac,C.,

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

COMMENT

Burkett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C., David, R., Delgado, O., Deshazo, D., Ding, Y., Domah-Rashid, N., Dugan-Rocha, S., Durbin, K.J., Fernandez, C., Ferraguto, D., Forcum-Tenney, J., Frantz, P., Ganesh, R., Gorrell, J.H., Gorrell, L.L., Guevara, W., Harris, K., Hernandez, J., Hodgson, A., Hogue, M., Holloway, C., Hosak, H., Jackson, L.E., Jackson, L., Jia, Y., Jones, M., Kelly, S., Kondejewski, N., Kong, Y., Kovar, C., Leal, B., Li, Z., Lichtarge, O., Liu, J., Liu, M., Logan, O., Lozano, R.J., Lu, J., Lucier, R., Martin, R., Martinez, C., McLeod, M.P., Mel, G., Morgan, M., Morris, S., Nash, S., Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S., Ouellet, M., Parish, B., Paxton, S., Payton, B., Perez, L., Pl, L.L., Quiles, M., Reiter, D., Rives, M., Samuel, S., Say, J., Scherer, S., Shah, B., Shen, H., Simon, M., Sparks, A., Stamps, A., Seng, R., Tabor, P., Taylor, T., Vasquez, L., Vinson, R., Vo, Q., Wahab, M., Wallington, S., Weinstein, G., Weinstein, I.R., Williamson, A., Morley, K., Wren, J., Wrenford, G., Yu, W., Zhou, X., Nelson, D. and Gbbs, R.

Direct Submission
Unpublished
2 (bases 1 to 169620)
Morley, K.C.

Direct Submission
Submitted (03-NOV-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Aug 7, 2000 this sequence version replaced gi:8705345.

----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HMOG
Center clone name: RP1-458H3
----- Summary Statistics
Assembly program: Phrap, version 0.990329
Consensus quality: 139025 bases at least Q40
Consensus quality: 154842 bases at least Q30
Consensus quality: 159725 bases at least Q20
Estimated insert size: 162720; sum-of-contris estimation
Estimated insert size: 171608; agarose-fp estimation
Quality coverage: 3.9x in Q20 bases; agarose-fp estimation
Quality coverage: 4.1x in Q20 bases; sum-of-contris estimation

NOTE: Estimated insert size may differ from sequence length
(see http://www.hgsc.bcm.tmc.edu/docs/genbank/draft_data.html).
NOTE: This is a 'working draft' sequence. It currently consists of 18 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 28689: contig of 28689 bp in length
28690 28789: gap of unknown length
28790 50832: gap of unknown length
50832 50932: gap of unknown length
50932 69144: contig of 18212 bp in length
69144 69244: gap of unknown length
69244 84204: contig of 14960 bp in length
84204 84304: gap of unknown length
84304 94667: contig of 10363 bp in length
94667 94767: gap of unknown length
94767 107261: contig of 12494 bp in length
107261 107361: gap of unknown length
107361 117550: contig of 10189 bp in length
117550 117650: gap of unknown length
117650 126939: contig of 9289 bp in length
126939 127039: gap of unknown length
127039 135040: contig of 8001 bp in length
135040 135140: gap of unknown length
135140 141639: contig of 6499 bp in length

141640 141739: gap of unknown length
141740 149558: contig of 7819 bp in length
149559 149658: gap of unknown length
149659 154562: contig of 4904 bp in length
154563 154662: gap of unknown length
154663 158987: contig of 4325 bp in length
158988 159087: gap of unknown length
159088 162377: contig of 3289 bp in length
162377 162476: gap of unknown length
162477 165191: contig of 2715 bp in length
165192 165291: gap of unknown length
165292 167173: contig of 1882 bp in length
167174 167273: gap of unknown length
167274 168393: contig of 1120 bp in length
168394 168493: gap of unknown length
168494 169620: contig of 1127 bp in length.

FEATURES
source
1..169620
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RP1-458H3"

BASE COUNT 52024 a 33180 c 32128 g 50322 t 1966 others
ORIGIN

Query Match 75.5% Score 299; DB 2: Length 169620;
Best Local Similarity 99.7% Pred. No. 3.2e-170;

Matches 349; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 ANGGGTGATCTTTTTCCTTCGAGGATCTTTTCATCTTTTCAGAGCACTTCGGGCG 60
DB 87441 ANGGGTGATCTTTTTCCTTCGAGGATCTTTTCATCTTTTCAGAGCACTTCGGGCG 87382
QY 61 GAGTATGTAAACTCTCTGGGTCTCTGTGTGCTGAGTGGCTGCTACTGAGACTCTG 120
DB 87381 GAGTATGTAAACTCTCTGGGTCTCTGTGTGCTGAGTGGCTGCTACTGAGACTCTG 87322
QY 121 CATACACAGCTCTGTATATACGAGCCAGGAGCCCTGTGTCATCTGGGCTACAGAGATCC 180
DB 87321 CATACACAGCTCTGTATATACGAGCCAGGAGCCCTGTGTCATCTGGGCTACAGAGATCC 87262
QY 181 CCTGATCTGTGGGTGCAAGATCTGTGGAGAGTGTGTTTCTCGATGGGCTACAC 240
DB 87261 CCTGATCTGTGGGTGCAAGATCTGTGGAGAGTGTGTTTCTCGATGGGCTACAC 87202
QY 241 AATCAGCTACCTCTCCCTGCTGGAGGTGGGCTTTTTCGTCATCTGCTGCC 300
DB 87201 AATCAGCTACCTCTCCCTGCTGGAGGTGGGCTTTTTCGTCATCTGCTGCC 87142
QY 301 TGGGGGGGAGTTGGCTACCCCATTTTTCATCTCTGTGGTCAAG 350
DB 87141 TGGGGGGGAGTTGGCTACCCCATTTTTCATCTCTGTGGTCAAG 87092

RESULT 2
AL137847/c 143372 bp DNA linear PRI 16-NOV-2001
LOCUS Human DNA sequence from clone RP11-439K3 on chromosome 9q22.2-31.1,
DEFINITION complete sequence.
ACCESSION AL137847
VERSION AL137847.12 GI:16973786
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens

REFERENCE 1 (bases 1 to 143372)
AUTHORS Kimberley, A.
TITLE Direct Submission
JOURNAL Submitted (16-NOV-2001) Wellcome Trust Sanger Institute, Hinxton,
Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
humquery@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk
On Nov 17, 2001 this sequence version replaced gi:16408610.

COMMENT


```

CC * 120299 122220: contig of 1922 bp in length
CC * 122321 122320: gap of unknown length
CC * 122321 123779: contig of 1459 bp in length.
XX
FH Key Location/Qualifiers
FH source 1. 123779
FT /chromosome="3"
FT /db_xref="taxon:9606"
FT /organism="Homo sapiens"
FT /clone="RP11-79K12"
XX
SQ Sequence 123779 BP: 37802 A; 24017 C; 23437 G; 37612 T; 911 other;

Query Match 10.1% Score 40; DB 30; Length 123779;
Best Local Similarity 100.0%; Pred. No. 2, le-12;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 69 AAAACTCTGGGCTCTGTGTGTGCTGAGTGGCTGCTCT 108
3083 AAAACTCTGGGCTCTGTGTGTGCTGAGTGGCTGCTCT 3044

RESULT 4
LOCUS AC076969 128118 bp DNA linear HTG 15-OCT-2001
DEFINITION Homo sapiens chromosome 3 clone RP11-79K12, WORKING DRAFT SEQUENCE,
AC076969
14 unordered pieces.
AC076969 6 GI:16117967
VERSION HTG; HTGS_PHASE1; HTGS_DRAFT.
KEYWORDS Homo sapiens.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 128118)
Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,
Aisbrook,S.L., Amaralungu,H.C., Are,J.R., Banks,T., Barbarta,J.,
Benton,U., Bimaga,K., Blankenburg,K., Bonnin,D., Bouck,J.,
Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C.,
Burck,P., Burckett,C., Burrell,K.L., Byrd,N.C., Carron,T.F.,
Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Cleveland,C.D., Cox,C.,
Chen,Z., Chowdhry,I., Christopoulos,C., Cleaveland,C.D., Chen,R.,
Coyle,M.D., Dalhorne,S.R., David,R., Davila,M.L., Davis,C.,
Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O.,
Dean,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H.,
Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C.,
Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J.,
Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T.,
Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S.,
Hamilton,K., Harris,C., Harris,K., Hart,M., Haylak,P., Hawes,A.,
Hernandez,J., Hernandez,O., Hodgson,A., Hogues,M., Holloway,C.,
Hollins,B., Homsli,F., Howard,S., Huber,J., Huylk,S., Hume,J.,
Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jollivet,S.,
Joudah,S., Karlsson,E., Kelly,S., Khan,U., King,L., Korvah,J.,
Kovar,C., Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C.,
Lewis,L., Li,U., Li,Z., Lichtarge,O., Lieu,C., Liu,C., Liu,W.,
Loussegod,H., Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R.,
Ma,J., Maheshwari,M., Mapua,P., Martin,R., Martindale,A.,
Martinez,E., Massey,E., Mawhinney,E., Mcleod,M.P., Meador,M.,
Mei,G., Metzker,M., Miner,G., Miner,D., Mitchell,T., Mohabbat,K.,
Morgan,M., Morris,S., Moser,M., Neal,Z., Newton,J., Newton,N.,
Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokwenwo,S.,
Ogun,M., Okunolu,G., Orazungu,N., Oviedo,R., Pace,A., Payton,B.,
Peery,M., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L.L.,
Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojibokan,I., Rolle,M.,
Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shooshitari,N.,
Stison,I., Sodergren,E., Sonaike,T., Sparks,A., Stanley,H.,
Stonon,A., Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K.,
Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N.,
Thomas,S., Usmani,K., Vasquez,L., Vera,V., Villalón,D., Vinson,R.,
Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C.,
Watlington,S., Williams,G., Williamson,A., Wleczek,R., Wooden,S.,

```

FEATURES

source

```

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT
Morley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,
Weinstock,G. and Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 128118)
Morley,K.C.
Submitted (01-NUC-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Oct 14, 2001 this sequence version replaced gi:10047573.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HBRJ
Center clone name: RP11-79K12
----- Summary Statistics
Sequencing vector: M13; L08821
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 111307 bases at least Q40
Consensus quality: 117905 bases at least Q30
Consensus quality: 121494 bases at least Q20
Estimated insert size: 122854; sum-of-contigs estimation
Quality coverage: 0x in Q20 bases; agarose-tp estimation
Quality coverage: 4x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/genbank/draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 14 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
1
20354: contig of 20354 bp in length
20355 20454: gap of unknown length
20455 33071: contig of 12617 bp in length
33072 33171: gap of unknown length
33172 46935: contig of 13764 bp in length
46936 47035: gap of unknown length
47036 57066: contig of 10031 bp in length
57067 57166: gap of unknown length
57167 70378: contig of 13212 bp in length
70379 70478: gap of unknown length
81786: contig of 11308 bp in length
81887 91428: gap of unknown length
91429 91528: contig of 9542 bp in length
91529 100453: gap of unknown length
100454 100555: contig of 8927 bp in length
100556 107879: gap of unknown length
107880 107979: contig of 7324 bp in length
107980 114149: gap of unknown length
114149 114249: contig of 6170 bp in length
114250 120186: gap of unknown length
120187 120286: gap of 5937 bp in length
120287 123236: contig of 2950 bp in length
123237 123336: gap of unknown length
123337 125606: contig of 2270 bp in length
125607 125706: gap of unknown length
125707 128118: contig of 2412 bp in length.
Location/Qualifiers
1. 128118
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RP11-79K12"

```


1042	1065	743	758	1204	1224
2783	2808	3045	2963	2722	2723
7074	7158	1288	1262	321	<800
4087	3861	967	973	1066	1058
7976	7926	461	<800	470	<800
1233	1199	4649	4618	3055	3072
3875	3861	5289	5417	10961	10848
27	<800	1098	1095	6725	6888
1137	1065	798	809	1896	1903
750	749	111	<800	8682	8666
214	<800	7644	7609	4438	4342
836	887	10084	9932	514	<800
1670	1651	3641	3589	3077	3072
2650	2609	7555	7609	1371	1337
1890	1872			4185	4087
4615	4522			719	<800
1321	1273			4130	4087
				367	<800
				2979	3072
				333	<800

TITLE	Homo sapiens chromosome 15, clone CTD-3049M7
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 148290)
AUTHORS	Birren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barina,N., Bastien,Y., Boguslavsky,L., Bouckgalter,B., Brown,A., Camarata,J., Campolano,A., Chang,J., Chazaro,B., Chopel,Y., Colangelo,M., Collins,S., Collamore,A., Cook,A., Cooke,P., Deatellano,K., Dewar,K., Diaz,J.S., Dodge,S., Ferrelita,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Girde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamel,A., Karats,A., Kells,C., Lavoque,K., Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G., Maclean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrum,J., Menus,L., Milhov,T., Mlenaga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollata,V., Raymond,C., Retta,R., Ribickad,M., Riley,R., Rise,C., Rogov,P., Roman,J., Roseltl,M., Roy,A., Santos,R., Schauer,S., Schupack,R., Seaman,S., Severy,P., Spencer,B., Strange-Thomann,N., Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Testay,S., Theodore,J., Topham,K., Travers,M., Travis,N., Triggillo,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W., Young,G., Zaimoun,J., Zembek,L., Zimmer,A. and Zody,M.
TITLE	Direct Submission
JOURNAL	Submitted (24-JAN-2002) Whitehead Institute/MIT center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
REFERENCE	3 (bases 1 to 148290)
AUTHORS	Birren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barina,N., Bastien,Y., Bloom,T., Boguslavsky,L., Bouckgalter,B., Brown,A., Camarata,J., Campolano,A., Chang,J., Chazaro,B., Chopel,Y., Colangelo,M., Collins,S., Collamore,A., Cook,A., Cooke,P., Deatellano,K., Dewar,K., Diaz,J.S., Dodge,S., Farro,S., Ferrelita,P., Fitzgerald,M., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Girde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamel,A., Karats,A., Kells,C., Lavoque,K., Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G., Maclean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrum,J., Menus,L., Milhov,T., Mlenaga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollata,V., Raymond,C., Retta,R., Ribickad,M., Riley,R., Rise,C., Rogov,P., Roman,J., Roseltl,M., Roy,A., Santos,R., Schauer,S., Schupack,R., Seaman,S., Severy,P., Spencer,B., Strange-Thomann,N., Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Testay,S., Theodore,J., Topham,K., Travers,M., Travis,N., Triggillo,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W., Young,G., Zaimoun,J., Zembek,L., Zimmer,A. and Zody,M.

```

FEATURES
SOURCE
    Location/Qualifiers
        1..128563
            /organism="Homo sapiens"
            /db_xref="taxon:9606"
            /chromosome="3"
            /clone="RP11-79K17"
            /clone_1fb="RPci human BAC library 11"
COUNT      37777 a 24752 c 24789 g 41265 t
IN
Query Match      10.1%; Score 40; DB 9; Length 128563;
Best Local Similarity 100.0%; Pred. No. 2.1e-12;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY      69 AAAAAGCTGCGGCTCTGCTGTGTGCGCTAGTGGCTGCTCT 108
        |||||||
DB       27283 AAAAAGCTGCGGCTCTGCTGTGTGCGCTAGTGGCTGCTCT 27244

```

RESULT 6				
AC107979/c				
LOCUS	AC107979	148290 bp	DNA	linear PRI 01-JUL-2002
DEFINITION	Homo sapiens chromosome 15, clone CTD-2049M7, complete sequence.			
ACCESSION	AC107979			
VERSION	AC107979.7	GI:21592043		
KEYWORDS	HTG.			
SOURCE	human.			
ORGANISM	Homo sapiens			
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;			
	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
REFERENCE	1 (bases 1 to 148290)			
AUTHORS	Birren,B., Nusbaum,C. and Lander,E.			

2 (bases 1 to 148290)

unpublished

Homo sapiens chromosome 15, clone CTD-3049M7

2 (bases 1 to 148290)

Birren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N.,
Anderson,S., Barra,N., Bastien,V., Boguslavsky,L., Boukhgalter,B.,
Brown,A., Camarata,J., Campolano,A., Chang,J., Chazaro,B.,
Chapel,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A.,
Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
Kamat,A., Karatas,A., Kells,C., Lacroque,K., Lamazates,R.,
Landers,T., Lehoczy,J., Levine,R., Liu,G., Maclean,C.,
Macdonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M.,
Mcwan,P., McKernan,K., Meldrim,J., Meneus,L., Mihova,T.,
Mienga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C.,
Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J.,
Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C.,
Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,
Rosetti,P., Roy,A., Santos,R., Schauer,S., Schnuppach,R., Seaman,S.,
Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission

Submitted (24-JAN-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 148290)

Birren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N.,
Anderson,S., Barra,N., Bastien,V., Bloom,T., Boguslavsky,L.,
Boukhgalter,B., Brown,A., Camarata,J., Campolano,A., Chang,J.,
Chazaro,B., Chapel,Y., Colangelo,M., Collins,S., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzgerald,M., Fitzhugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I.,
Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Lacroque,K.,
Lamazates,R., Landers,T., Lehoczy,J., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Marquis,N.,
Matthews,C., McCarthy,M., McEwan,P., McKernan,K., Meldrim,J.,
Meneus,L., Mihova,T., Mienga,V., Murphy,T., Naylor,J., Nguyen,C.,
Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C.,
Rogov,P., Roman,J., Rosetti,P., Roy,A., Santos,R., Schauer,S.,
Schnuppach,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Testaye,S.,
Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J.,
Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission

Submitted (26-JUN-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
4 (bases 1 to 148290)

Birren,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S.,
Barra,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhgalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepley,Y., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mienga,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schnuppach,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.

TITLE Direct Submission
JOURNAL Submitted (01-JUL-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jun 26, 2002 this sequence version replaced gi:21321840.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WITR

Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information
Center project name: L24533
Center clone name: 3049_M_7

FEATURES

Source Location/Qualifiers
1. 148290
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15"
/clone="CTD-3049M7"
/clone_id="CTD2 Human BAC"
complement(1982..2092)
repeat_region
/rpt_family="MIR"
complement(3356..3654)
repeat_region
/rpt_family="AluSx"
3789..3844
/rpt_family="MER58"
complement(4429..4728)
repeat_region
/rpt_family="AluSp"
5494..5723
repeat_region
/rpt_family="(TA)n"
5727..5766
repeat_region
/rpt_family="(GA)n"
5806..5857
repeat_region
/rpt_family="AT-rich"
6377..6581
repeat_region
/rpt_family="L1"
6647..7069
repeat_region
/rpt_family="L1ME3"
7057..7631
repeat_region
/rpt_family="L1ME3"
complement(8821..8958)
repeat_region
/rpt_family="MIR3"
11129..11224
repeat_region
/rpt_family="FRAM"
11129..11338
repeat_region
/rpt_family="CT-rich"
11943..12090
repeat_region
/rpt_family="MIR"
12818..13060
repeat_region
/rpt_family="MIR"
13118..19146
repeat_region
/rpt_family="L1PA2"
20846..20912
repeat_region
/rpt_family="CT-rich"
21294..21314
repeat_region
/rpt_family="AT-rich"
21635..23417
repeat_region
/rpt_family="L1PA16"
23438..25940
repeat_region
/rpt_family="L1PA16"
26121..26145
repeat_region
/rpt_family="(TTCA)n"
26293..26417
repeat_region
/rpt_family="MIR"
complement(26872..26937)
repeat_region
/rpt_family="L2"
complement(27064..27431)
repeat_region
/rpt_family="L2"
27917..28357

repeat_region /rpt_family="MLT1B"
28460..28806
repeat_region /rpt_family="MLT1I"
29509..29698
repeat_region /rpt_family="MIR"
29716..29790
repeat_region /rpt_family="CT-rich"
29791..29811
repeat_region /rpt_family="AT-rich"
complement(31182..31787)
repeat_region /rpt_family="L2"
complement(32398..32581)
repeat_region /rpt_family="AluSg"
complement(32582..32777)
repeat_region /rpt_family="L1PB1"
complement(32778..32912)
repeat_region /rpt_family="AluSg"
33104..33207
repeat_region /rpt_family="L1MC5"
complement(33208..33365)
repeat_region /rpt_family="MER53"
33366..33529
repeat_region /rpt_family="L1MC5"
33877..34506
repeat_region /rpt_family="L1MED"
34667..34808
repeat_region /rpt_family="L1MED"
35585..36698
repeat_region /rpt_family="L1M4"
complement(36699..36830)
repeat_region /rpt_family="FLAM_C"
36831..38141
repeat_region /rpt_family="L1M4"
38194..38373
repeat_region /rpt_family="MLT1I"
39530..39757
repeat_region /rpt_family="MIR"
39992..40017
repeat_region /rpt_family="(TTTTG)n"
40283..40583
repeat_region /rpt_family="AluY"
41414..41616
repeat_region /rpt_family="MLT1C"

Query Match 10.1%; Score 40; DB 9; Length 148290;
Best Local Similarity 100.0%; Pred. No. 2.1e-12;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 68 TAAACTCTGGGCTCTGTGTGCTGAGTGCGTCTC 107
Db 84499 TAAACCTCTGGGCTCTGTGTGCTGAGTGCGTCTC 84460

RESULT 7
AC103996/c 165649 bp DNA linear PRI 01-JUL-2002
LOCUS
DEFINITION Homo sapiens chromosome 15, clone RP11-76E17, complete sequence.
AC103996
VERSION AC103996.7 GI:21637504
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 165649)
REFERENCE
AUTHORS Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N.,
Anderson, S., Barina, N., Bastien, V., Boguslavskiy, L., Boukhgalter, B.,
Brown, A., Camarata, J., Campio, A., Chang, J., Chazaro, B.,
Choepel, Y., Colangelo, M., Collins, S., Collymore, A., Cook, A.,
JOURNAL 2 (bases 1 to 165649)
REFERENCE
AUTHORS
TITILE
JOURNAL
REFERENCE
AUTHORS

TITLE
JOURNAL
REFERENCE
AUTHORS

Cooke, P., DeArfollano, R., Dewar, K., Diaz, J.S., Dodde, S., Fero, S., Ferrelia, P., Fitzhugh, M., Gage, D., Galgan, J., Gardyna, S., Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N., Hages, B., Heatford, A., Horton, L., Hulme, M., Iliev, I., Jones, C., Kamat, A., Karatas, A., Kells, C., Lacroque, K., Macleates, R., Landers, T., Lenocksky, J., Levine, R., Liu, G., Maclean, C., Macdonald, P., Major, J., Margus, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., McPheeters, R., Meldrum, J., Meneses, L., Mihova, T., Mlekena, V., Murphy, T., Naylor, J., Nguyen, C., Norby, C., Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunhphan, P., Pierre, N., Pollara, V., Raymond, C., Retta, R., Ribback, M., Riley, R., Rise, C., Rodov, P., Roman, J., Roselli, M., Roy, A., Santos, R., Schauer, S., Schyback, R., Strauss, S., Severly, P., Spencer, B., Strange-Thomann, N., Stojanovic, N., Stearns, N., Subramanian, A., Talamas, J., Teefaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Triggillo, J., Vassiliou, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.-J., Young, G., Zaitoun, J., Zembek, D., Zimmer, A. and Zody, M.

Direct Submission

Submitted (01-DEC-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 165649)

(bases 1 to 165649)

Barren, B., Busbaum, C., Lander, E. A. I., Allen, N., Anderson, S., Barria, N., Bastien, V., Bloom, T., Boussilavsky, L., Bourkhalter, B., Cameraret, J., Chang, J., Chazaro, B., Choepeli, Y., Collamore, A., Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Fario, S., Ferrello, P., Fitzgerald, M., Gage, D., Galagan, J., Garryna, S., Gird, S., Graham, L., Grand-Pierre, N., Haags, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kanat, A., Karlas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K., Liu, G., Maclean, C., Macdonald, P., Major, J., Matthews, C., McCarthy, M., Meldrum, J., Meunus, L., Milnova, T., Mieng, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhuan, P., Plerre, N., Raymond, C., Retta, R., Risse, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schnuppach, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Stange, Thomann, N., Stojanovic, N., Talamas, J., Testaye, S., Theodore, J., Topham, K., Travers, M., Vassiliou, H., Vieler, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zimmer, L., Zimmer, A. and Zody, M.

Direct Submission

Submitted (26-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

Research, 320 Charles Street, Cambridge, MA 02141, USA

4 (bases 1 to 16549)

Barren, B., Nusbbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
Barnett, N., Bastien, V., Bloom, T., Boguslavsky, L., Boukhalter, B.,
Cammarata, J., Chang, J., Chazaro, B., Choepel, Y., Collymore, A.,
Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S.,
Faro, S., Ferreira, P., Fitzgerald, M., Gage, D., Galagan, J.,
Gardys, S., Gord, S., Graham, L., Grand-Pierre, N., Haas, B.,
Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Katat, A.,
Karatas, A., Kellis, C., Landers, T., Levine, R., Lindblad-Toh, K.,
Liu, G., Maclean, C., Macdonald, P., Major, J., Matthews, C.,
McCarthy, M., Meldrum, J., Menous, L., Mihov, T., Miemala, V.,
Murray, T., Naylor, J., Nguyen, C., Nicot, R., Norbu, C., Norman, C. H.,
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
Phanangkarn, P., Pierre, N., Raymond, C., Rella, R., Rise, C., Rogov, P.,
Roman, J., Roy, A., Schauer, S., Schupbach, R., Seaman, S., Severy, P.,
Smith, C., Spencer, B., Stange-Riommann, N., Stojanovic, N., Talamas, J.,
Teisfaye, S., Theodore, J., Totham, K., Travers, M., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.

Direct Submission

Submitted (01-JUL-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

On Jul 1, 2002 this sequence replaced gi:21592191.

All repeats were identified using RepeatMasker:

SMILE, A.F.F.A. & Green, P. (1996-1997)

<http://ltp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBH

Web site: <http://www-seq.wi.mit.edu>

FEATURES

Contact: sequence_submissions@genome.wi.mit.edu

Project Information
Center project name: L21917
Center clone name: 76_E_17

URES	Location/Qualifiers
source	1. 165649 /organism="Homo sapiens" /db_xref="taxon:9606" /chromosome="15" /map="15" /clone="PPI1-76E17" /clone.lib="RPCT-11 Human Male BAC complement(2. 865) /rpt_family="L1PA13" complement(664. .894) /rpt_family="L1PA13" complement(895. .1248) /rpt_family="THE1A" complement(1249. .1504) /rpt_family="L1PA13" 1505. .1626 /rpt_family="AluXs" 1627. .1659 /rpt_family="CMAA)n" 1660. .1831 /rpt_family="AluXs" complement(1832. .3132) /rpt_family="L1PA13" complement(3140. .3699) /rpt_family="L1MC4c complement(4461. .4755) /rpt_family="AluXs" 5027. .5397 /rpt_family="L2" complement(5650. .6330) /rpt_family="L1ME1" complement(6345. .6460) /rpt_family="L1ME1" 7214. .7319 /rpt_family="L1MC3" 7354. .7655 /rpt_family="L1MC3" 7659. .7743 /rpt_family="L1PA10" complement(7745. .7843) /rpt_family="AluSp/q" 7844. .14156 /rpt_family="L1PA10" 14198. .14328 /rpt_family="AluY" 14340. .14503 /rpt_family="T(A)n" 14507. .15032 /rpt_family="L1MC3" 15036. .15320 /rpt_family="L1MC3" complement(15218. .16449) /rpt_family="L1PA4" 16450. .18154 /rpt_family="L1PA4" 18155. .19310 /rpt_family="L1MC3" 19211. .19266 /rpt_family="T(A)n" 19267. .19372 /rpt_family="L1MC3" 19401. .19476 /rpt_family="T(TATA)n" 19483. .19546 /rpt_family="C(CATATA)n" 19566. .19625 /rpt_family="AT-rich"

```

repeat_region      19696..19750
/rpt_family="GA-rich"
repeat_region      19752..19920
/rpt_family="L1MD3"
repeat_region      19994..20102
/rpt_family="L2"
unsure             20049..20114
/note="single clone coverage"
repeat_region      20485..20655
/rpt_family="MIR3"
repeat_region      complement(21285..21441)
/rpt_family="MIR"
repeat_region      21496..21717
/rpt_family="L2"
repeat_region      complement(21720..21796)
/rpt_family="MIR"
repeat_region      complement(21943..22131)
/rpt_family="MIR"
repeat_region      23082..23195
/rpt_family="L2"
repeat_region      23198..23248
/rpt_family="GA-rich"
repeat_region      23267..23333
/rpt_family="CATR)n"
repeat_region      complement(24465..24833)
/rpt_family="MTT1A2"
repeat_region      26142..26334
/rpt_family="MIR"
repeat_region      complement(27659..27811)
/rpt_family="MIR"
repeat_region      complement(28811..28880)
/rpt_family="MIR"
repeat_region      complement(29026..29269)
/rpt_family="MIR"
repeat_region      complement(29673..29706)
/rpt_family="MSTB"

Query Match      10.1%, Score 40; DB 9; Length 165649;
Best Local Similarity 100.0%; Pred. No. 2.1e-12;
Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 68 TAAACTCTGGGCTCTGTGTGTCCTGAGTGGCTGCTC 107
|||||
Db 8110 TAAACTCTGGGCTCTGTGTGTCCTGAGTGGCTGCTC 8071

RESULT 8
AC104303 178650 bp DNA linear PRI 25-FEB-2002
AC104303
AC104303 AC064830
AC104303.2 GI:18874945
HTG.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 178650)
AUTHORS Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Saenphimachak,C., Phelps,K.A., Raymond,C. and Haugen,E.D.
TITLE Direct Submission
JOURNAL Unpublished
AUTHORS 2 (bases 1 to 178650)
Kaul,R.K., Olson,M.V., Raymond,C. and Haugen,E.D.
TITLE Direct Submission
JOURNAL Submitted (07-DEC-2001) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 178650)
AUTHORS Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Saenphimachak,C., Phelps,K.A., Raymond,C. and Haugen,E.D.
TITLE Direct Submission
JOURNAL Submitted (25-FEB-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA

```

COMMENT

On Feb 25, 2002 this sequence version replaced gi:17402782.

----- Genome Center

Center: University of Washington Genome Center

Center Code: UMG

Web site: <http://www.genome.washington.edu>

Contact: wgchits@u.washington.edu

Drafting Center: WUGSC

----- Project Information

Center project name: chr-3

Genet clone name: RP11-391P4 (bc0402)

----- Summary statistics

Sequencing vector: unknown; 55% of reads

Sequencing vector: plasmid; 45% of reads

Chemistry: Dye-terminator ET; 89% of reads

Chemistry: Dye-terminator Big Dye; 11% of reads

Assembly program: Phrap; version 0.990319

Consensus quality: 178494 bases at least 940

Consensus quality: 178650 bases at least 920

Insert size: 178648; sum-of-contigs

Quality coverage: 8.0x in Q20 bases; sum-of-contigs

Overlapping Sequences:

5': RP11-475023 (UMGC:bc0439) AC023346

3': RP11-79K12 AC076969

Sequence Quality Assessment:

This entry has been annotated with sequence quality estimates computed by the Phrap assembly program.

All manually edited bases have been reduced to quality zero.

Quality levels above 40 are expected to have less than

1 error in 10,000 bp.

Base-by-base quality values are not generally visible from the

Genbank flat file format but are available as part

of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted:

all regions were either double-stranded or sequenced with an

alternate chemistry or covered by high quality data (i.e., Phred

quality >= 30); an attempt was made to resolve all sequencing

problems, such as compressions and repeats; all regions were

covered by at least one plasmid subclone or more than one M13

subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest

fingerprinting. Comparison of the experimentally derived digest

fragments with sequence-predicted fragments is given below.

The electronically-digested sequence consists of both insert and

vector, in order to accurately represent the entire circular BAC.

Small fragments below a variable cutoff (approximately 400-800 bp)

are not resolved in the fingerprint and hence do not appear

in the table. There are no significant remaining discrepancies

between the experimental and predicted values. Uniquely ordered

fragments are separated by dashed lines.

BgIII

EcORI

HindIII

SeqDerMap FngPrnt SeqDerMap FngPrnt SeqDerMap FngPrnt

7050 7417 8696 8715 7163 7098

2067 2138 6 800 6382 6518

13472 13264 3077 3083 512 800

7644 7859 514 800 449 800

111 800 4438 4377 7988 7884

798 783 8682 8715 11779 11698

1098	1109	1896	1876	472	<800
5288	5395	6724	6948	783	787
4649	4507	10961	10773	926	926
461	<800	3055	3083	2431	2538
967	982	470	<800	1015	1029
1288	1268	1066	1037	758	<800
3045	2923	321	<800	1550	1541
743	783	2722	2683	361	<800
5628	5682	1204	1300	763	<800
2714	2699	6992	7281	2975	3002
4002	3900	7210	7830	3299	3468
227	<800	7821	8101	1907	1894
1794	1784	6855	6948	4020	4220
2168	2138	859	857	5158	5126
10032	9744	2629	2683	497	<800
6694	6810	1065	1037	4321	4220
1505	1491	3934	3965	887	926
3312	3304	859	857	180	<800
896	912	931	935	4699	4887
5873	5938	858	857	4755	4674
78	<800	799	857	26	<800
2135	2138	1895	1876	906	926
133	<800	58	<800	2325	2346
11939	11575	5129	5076	369	<800
1507	1491	6051	6048	2523	2538
8069	8181	3538	3537	1889	1894
1632	1617	1114	1094	4437	4428
3729	3678	8420	8391	373	<800
5533	5395	1024	1037	24	<800
358	<800	1323	1380	263	<800
3764	3678	1502	1481	632	<800
7022	6810	4538	4539	1258	1247
8534	8821	1414	1380	1114	1083
5206	5050	521	<800	1370	1320
9052	9744	8346	8391	1262	1247
1625	1617	39	<800	1546	1541
749	783	2716	2683	459	<800

Query Match 10.1%; Score 40; DB 9; Length 178650;
 Best Local Similarity 100.0%; Pred. No. 2, 1e-12;
 Matches 40; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 69 AAACTCCTGGCTCTGTGTGCTGAGTGGCTGCTCT 108
 Db 165449 AAAACTCCTGGCTCTGTGTGCTGAGTGGCTGCTCT 165488

RESULT 9
 AC090762/c 192826 bp DNA linear PRI 28-FEB-2002
 LOCUS Homo sapiens chromosome 15, clone RP11-387E8, complete sequence.
 DEFINITION AC090762
 AC090762.9 GI:18997378
 VERSION
 KEYWORDS
 SOURCE
 ORGANISM
 Homo sapiens.
 Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE
 1 (bases 1 to 192826)
 Birren, B., Linton, L., Nusbaum, C. and Lander, E.
 Unpublished
 2 (bases 1 to 192826)
 Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S.,
 Barne, N., Bastien, V., Boguslavsky, L., Boukhalter, B., Brown, A.,
 Camarata, J., Campopiano, A., Choepel, Y., Colangelo, M., Collins, S.,
 Collamore, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S.,
 Dodge, S., Faro, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J.,
 Gargay, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N.,
 Hagos, B., Heaford, A., Horton, L., Hulme, M., Iliev, I., Johnson, R.,
 Jones, C., Karatas, A., Larocque, K., Lamazares, R., Landers, T.,
 Lehocck, J., Levine, R., Liu, G., Maclean, C., Macdonald, P.,
 Marquis, N., Mathews, C., McCarthy, M., McEwan, P., McKernan, K.,
 McPheters, R., Meldrim, J., Menus, L., Mihova, T., Mienga, V.,
 Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C. H.,
 O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
 Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Retta, R.,
 Ribback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M.,
 Roy, A., Santos, R., Schauer, S., Schnupack, R., Seaman, S., Severy, P.,
 Sougnaz, C., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
 Strauss, N., Subramanian, A., Talamas, J., Testaf, S., Theodore, J.,

Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE
Submitted (10-MAR-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

JOURNAL
3 (bases 1 to 192826)

REFERENCE
Birren, B., Linton, L., Nusbaum, C., Lander, E., All, A., Allen, N., Anderson, S., Barna, N., Bastien, V., Boguslavsky, L., Boukhalter, B., Brown, A., Camarata, J., Campopiano, A., Chang, J., Chazaro, B., Choquel, Y., Collangelo, M., Collins, S., Collamore, A., Cook, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J. S., Dodge, S., Faro, S., Ferreira, P., Fitzhugh, W., Gage, D., Galagan, J., Gardyna, S., Ginde, S., Gord, S., Goyette, M., Graham, L., Grand-Pierre, N., Hago, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A., Karatas, A., Kells, C., Larocque, K., Lamazares, R., Landers, T., Lebeck, J., Levine, R., Liu, G., Maclean, C., Macdonald, P., Major, J., Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K., Meldrum, J., Meneses, L., Mihova, T., Mlenga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C. H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K., Phunkhang, P., Pierre, N., Pollara, V., Raymond, C., Rella, R., Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rosetti, M., Roy, A., Santos, R., Schauer, S., Schupbach, R., Seaman, S., Severy, P., Spencer, B., Stange-Thomann, N., Stojanovic, N., Strauss, N., Sudrmanian, A., Talamas, J., Testaye, S., Theodore, J., Topham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W. J., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE
Submitted (28-FEB-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA

JOURNAL
On Feb 28, 2002 this sequence version replaced g1.18377189.

COMMENT
All repeats were identified using RepeatMasker:
Smith, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RN/RepeatMasker.html>

FEATURES
source

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www.seq.wi.mit.edu>
Contact: sequence.submissions@genome.wi.mit.edu

Project Information
Center Project name: L12392
Center Clone name: 367_E_8

Location/Qualifiers
1. 192826
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15"
/clone="RP11-387E8"
/clone_lib="RPC1-11 Human Male BAC"
complement(864..1222)
/rpt_family="L1PB3"
1223..1252
/rpt_family="(TAGG)n"
complement(1253..1492)
/rpt_family="L1PB3"
4287..4576
/rpt_family="AluDb"
4671..5034
/rpt_family="THE1B"
5130..5438
/rpt_family="AluSx"
5517..5662
/rpt_family="MIR"
5798..5827
/rpt_family="AT_rich"
5985..6114
/rpt_family="AluDb"
6227..6253
/rpt_family="(CA)n"

repeat_region 7329..7446
/rpt_family="MIR"
complement(8147..8452)
repeat_region 8540..8639
/rpt_family="AluDb"
complement(8632..8916)
repeat_region 10391..10599
/rpt_family="L2"
repeat_region 11688..11828
/rpt_family="MER3"
12029..12078
/rpt_family="AT_rich"
12092..12447
/rpt_family="THE1C"
complement(13616..13751)
repeat_region 14142
/rpt_family="MIR"
complement(15277..15553)
repeat_region 16005
/rpt_family="MER8"
complement(16101..16440)
repeat_region 16920..16958
/rpt_family="L3"
complement(17145..17444)
repeat_region 19953
/rpt_family="L1MEC"
complement(19978..20262)
repeat_region 20794
/rpt_family="L1MEC"
complement(20822..21097)
repeat_region 21743
/rpt_family="L1MEC"
22599..23518
/rpt_family="L1MEC"
23527..23901
/rpt_family="L1MEC"
repeat_region 24026
/rpt_family="MSTB1"
complement(24027..24256)
repeat_region 24557
/rpt_family="MER30"
complement(24257..24557)
repeat_region 24563..24594
/rpt_family="MSTB1"
24594
/rpt_family="AT_rich"
complement(25258..25639)
repeat_region 25638
/rpt_family="L1MCC"
26167..26506
/rpt_family="THE1B"
26746..27094
/rpt_family="Tiger2a"
27095..27184
/rpt_family="MADE1"
27185..27279
/rpt_family="Tiger2a"
28321..28363
/rpt_family="(TG)n"
29172..29333
/rpt_family="MIR"
30664..30898
/rpt_family="L1MB8"
31577..31598
/rpt_family="AT_rich"
32378..32475

```
repeat_region      /rpl_family="CT-rich"
                    34655..34960
                    /rpl_family="AluY"
repeat_region      complement(35296..35389)
                    /rpl_family="MER5B"
repeat_region      complement(35419..35517)
                    /rpl_family="L1MC4"
repeat_region      complement(35518..35823)
                    /rpl_family="AluSx"
repeat_region      complement(35824..36030)
                    /rpl_family="L1MC4"
repeat_region      36682..36758
                    /rpl_family="MER5A"
repeat_region      36975..37160
                    /rpl_family="MER5A"
repeat_region      38140..38353
                    /rpl_family="MER96B"
repeat_region      38633..38656
                    /rpl_family="(TTTA)n"
repeat_region      complement(38657..40056)
                    /rpl_family="L1PAS"
repeat_region      complement(40060..40250)
```

Query Match	10.1%	Score 40;	DB 9;	Length 192826;
Best Local Similarity	100.0%	Pred. No. 2.1e-12;		
Matches 40;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0

Qy 68 TAAACTCCTGGCTCTGTGTGTGCTTGAATGGCTGCTC 107
 |||||
 Db 190946 TAAACTCCTGGCTCTGTGTGTGCTTGAATGGCTGCTC 190907

RESULT 10	AC007445	32918 bp	DNA	linear	HTG 30-JUN-2000
LOCUS	AC007445				
DEFINITION	Homo sapiens chromosome 18 clone RP11-34A7 map 18,				*** SEQUENCING
	IN PROGRESS ***	1	Ordered piece.		
ACCESSION	AC007445				
VERSION	AC007445.5	GI:8844149			
KEYWORDS	HTGS_PHASE2.				
SOURCE	Homo sapiens.				
ORGANISM	Homo sapiens				

REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS

Eukaryota: Metazoa: Chordata: Craniata: Vertebrata: Euteleostomi: Mammalia: Eutheria: Primates: Catarrhini: Homnidae: Homo.
1 (bases 1 to 32918)
Birren, B., Linton, L., Nusbaum, C. and Lander, E.
Homo sapiens chromosome 18, clone RP11-344B7
Unpublished
2 (bases 1 to 32918)
Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, J.,

DAVEY, V., BALWIND, J., BARNIA, N., BECKERLEY, R., BERNH, J., BROWN, A., CASTLE, A., CERNY, J., COLANGELO, M., COLLINS, S., COLLYMORE, A., COOKE, P., DEARELLANO, K., DEPARRE, E., DEVON, K., DEWAR, K., DONELAN, L., DOYLE, M., FERREIRA, P., FITZHUGH, M., FORREST, C., FUNKE, R., GAGE, D., GALAGAN, J., GARDYNA, S., GILBERT, D., GRANT, G., HAGOS, B., HEALDOR, A., HORTON, L., HOWLAND, J. C., JONES, C., KANN, L., KARTAS, A., LEHOCZKY, J., LIEN, C., LOCKE, K., MACDONALD, P., MARDIN, N., MCGEAN, P., MCGURK, M., MCKERNAN, K., MCLAUGHLIN, J., MELDRUM, J., MOLL, M., MORRIS, W., MOWEN, J., MYCHALECKY, T., NAYLOR, J., NILIOFF, M., O'CONNOR, T., O'DONNELL, P., PAVLIN, B., PETERSON, K., POLLARA, V., RILEY, R., ROBERTS, D., ROY, A., SEVERY, P., STANGE-THOMANN, N., STOJANOVIC, N., STONE, C., SUBRAMANIAN, A., TESTAË, S., TORRELLA-MILLER, I., VASSILIEV, H., VO, A., WAGNER, A., WHEELER, J., WU, X., WYMAN, D., YE, W. J. and ZODY, M.

TITLE	Direct Submission
JOURNAL	Submitted (30-APR-1999) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
COMMENT	On Jun 30, 2000 this sequence version replaced gi:8705092.

Smith, A.F.A. & Green, P. (1990-1991)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: MIBR
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence.submissions@genome.wi.mit.edu

Project Information
Center project name: L571
Center clone name: 344_B-7

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 1 contigs. Gaps between the contigs
 * are represented as runs of N. The order of the pieces
 * is believed to be correct as given, however the sizes
 * of the gaps between them are based on estimates that have
 * been provided by the submitter.
 * This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.
 * 1 32918: contig of 32918 bp in length.

FEATURES	Location/Qualifiers
source	1. .32918

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="18"
/map="18"
/clone="RP11-344B7"
/clone_1b="RPCC1.11 Human Male BAC"
BASE COUNT      9462 a   6493 c   6065 g   9853 t   245 others
ORIGIN
```

Query Match	9.8%;	Score 39;	DB 2;	Length 32918;
Best Local Similarity	100.0%;	Pred. No. 8.2e-12;		
Matches 39;	Conservative 0;	Mismatches 0;	Indels 0;	Gaps 0;

Qy 60 GGAGTATGTAACCTCGTGGGTCTCTGTGTGTCCTGAG 98
|||||
Db 3535 GGAGTATGTAACCTCGTGGGTCTCTGTGTGTCCTGAG 3573

RESULT 11	
AL358817	
LOCUS	38936 bp
DEFINITION	Human DNA sequence from clone RP11-399N22 on chromosome 10,
	complete sequence.
ACCESSION	AL358817
VERSION	AL358817.18
KEYWORDS	GI:15990637
	HTG.
SOURCE	human.
ORGANISM	Homo sapiens

REFERENCE AUTHORS JOURNAL TITLE	COMMENT
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo. 1 (bases 1 to 38936) Lovell, J.	
Submitted (06-OCT-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humbureys@sanger.ac.uk requests: cloneenquiries@sanger.ac.uk On Oct 9, 2001 this sequence version replaced gi:14669268.	

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences without a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Emn., EMBL, Sv., SWISSPROT, Tr., TrEMBL, Wp., WormPeP; information on the WormPeP database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 10, constructed by the Sanger Centre Chromosome 10 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/MGP/chr10>

RP11-399N22 is from the library RPCI-11.2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/dacpac/home.htm>

VECTOR: pBAC3.6

IMPORTANT: This sequence is not the entire insert of clone RP11-399N22. It may be shorter because we sequence overlapping sections only once, except for a short overlap. The true left end of clone RP11-432J9 is at 36937 in this sequence. The true right end of clone RP11-91A1 is at 2000 in this sequence.

FEATURES

source
1. 38936
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="10"
/clone="RP11-399N22"
/clone_1lb="RPCI-11.2"
9315 a 9079 c 9111 g 11431 t
ORIGIN

Query Match 9.8%; Score 39; DB 9; Length 38936;
Best Local Similarity 100.0%; Pred. No. 8.2e-12;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Caps 0;

QY 60 GGAGTATGTAAACCTCGGTCTCTGTGCTGCTGAG 98
|||||
Db 28896 GGAGTATGTAAACCTCGGTCTCTGTGCTGCTGAG 28934

RESULT 12
LOCUS AC025179 124271 bp DNA linear HTG 20-APR-2001
DEFINITION Homo sapiens chromosome 5 clone CTD-2174B5, WORKING DRAFT SEQUENCE,
8 unordered pieces.
AC025179
AC025179.4 GI:13699647
HTG: HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEPIN.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 124271)
DOE Joint Genome Institute.
Sequencing of Human Chromosome 5
Unpublished
2 (bases 1 to 124271)
DOE Joint Genome Institute.
Direct Submission
Submitted (07-MAR-2000) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
On Apr 20, 2001 this sequence version replaced gi:7711794.
-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: <http://www.jgi.doe.gov>

Project Information
Center Project Name: 694394
Center clone name: CITB-HL_2174B5

Summary Statistics
Consensus quality: 116609 bases at least Q40
Consensus quality: 120226 bases at least Q30
Consensus quality: 121274 bases at least Q20
Estimated insert size: 117160; agarose-fp estimation
Estimated insert size: 123571; sum-of-contigs estimation
Quality coverage: 6.46 in Q20 bases; agarose-fp estimation
Quality coverage: 6.12 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently

* consists of 8 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1 1052: contig of 1052 bp in length
* 1053 1152: gap of unknown length
* 1153 3331: contig of 2179 bp in length
* 3332 3431: gap of unknown length
* 3432 7370: contig of 3939 bp in length
* 7371 7470: gap of unknown length
* 7471 15483: contig of 8013 bp in length
* 15484 15583: gap of unknown length
* 15584 24916: contig of 9333 bp in length
* 24917 25016: gap of unknown length
* 25017 39922: contig of 14506 bp in length
* 39923 40022: gap of unknown length
* 40023 68684: contig of 28662 bp in length
* 68685 68784: gap of unknown length
* 68785 124271: contig of 55487 bp in length.
Location/Qualifiers
1. 124271
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="5"
/clone="CTD-2174B5"
/clone_1lb="Caltech human BAC library D"

BASE COUNT 36863 a 24233 c 23599 g 38876 t 700 others
ORIGIN

Query Match 9.8%; Score 39; DB 2; Length 124271;
Best Local Similarity 100.0%; Pred. No. 8.6e-12;
Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAAACCTCGGTCTCTGTGCTGCTGAGT 99
|||||
Db 79646 GAGTATGTAAACCTCGGTCTCTGTGCTGCTGAGT 79684

RESULT 13
LOCUS AC008814 146671 bp DNA linear PRI 31-OCT-2001
DEFINITION Homo sapiens chromosome 5 clone CTD-217L12, complete sequence.
AC008814
AC008814.6 GI:16554342
HTG:
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 146671)
DOE Joint Genome Institute and Stanford Human Genome Center.
Direct Submission
Submitted (03-AUG-1999) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
On Oct 31, 2001 this sequence version replaced gi:15290309.
Draft Sequence Produced by DOE Joint Genome Institute
www.jgi.doe.gov

Project Information
Center Project Name: 694394
Center clone name: CITB-HL_2174B5

Summary Statistics
Consensus quality: 116609 bases at least Q40
Consensus quality: 120226 bases at least Q30
Consensus quality: 121274 bases at least Q20
Estimated insert size: 117160; agarose-fp estimation
Estimated insert size: 123571; sum-of-contigs estimation
Quality coverage: 6.46 in Q20 bases; agarose-fp estimation
Quality coverage: 6.12 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently

Project Information
Center Project Name: 694394
Center clone name: CITB-HL_2174B5

Summary Statistics
Consensus quality: 116609 bases at least Q40
Consensus quality: 120226 bases at least Q30
Consensus quality: 121274 bases at least Q20
Estimated insert size: 117160; agarose-fp estimation
Estimated insert size: 123571; sum-of-contigs estimation
Quality coverage: 6.46 in Q20 bases; agarose-fp estimation
Quality coverage: 6.12 in Q20 bases; sum-of-contigs estimation.
* NOTE: This is a 'working draft' sequence. It currently

* 153555 156134: contig of 2580 bp in length
* 156135 156234: gap of 100 bp
* 156235 158089: contig of 1855 bp in length
* 158090 158189: gap of 100 bp
* 158190 159747: contig of 1558 bp in length.

FEATURES

source

1. 159747

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="18"

/map="18p11.3"

/clone="RP11-752111"

1. 18444

/note="assembly_fragment"

18545. .29631

/note="assembly_fragment"

29732. .43143

/note="assembly_fragment"

43244. .57170

/note="assembly_fragment clone_end:SP6 vector_side:left"

57271. .68240

/note="assembly_fragment"

68341. .78134

/note="assembly_fragment"

78235. .87873

/note="assembly_fragment"

87974. .96197

/note="assembly_fragment"

96298. .103976

/note="assembly_fragment clone_end:T7 vector_side:left"

104077. .110726

/note="assembly_fragment"

110827. .116866

/note="assembly_fragment"

116967. .122789

/note="assembly_fragment"

122890. .127665

/note="assembly_fragment"

127766. .132618

/note="assembly_fragment"

132719. .136051

/note="assembly_fragment"

136132. .139749

/note="assembly_fragment"

139850. .142810

/note="assembly_fragment"

142911. .145855

/note="assembly_fragment"

145956. .148840

/note="assembly_fragment"

148941. .151243

/note="assembly_fragment"

151344. .153454

/note="assembly_fragment"

153555. .156134

/note="assembly_fragment"

156235. .158089

/note="assembly_fragment"

158190. .159747

/note="assembly_fragment"

BASE COUNT 47302 a 30397 c 29897 g 49849 t 2302 others
ORIGIN

Query Match 9.8%; Score 39; DB 2; Length 159747;
Best Local Similarity 100.0%; Pred. No. 8.6e-12;

Matches 39; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 60 GGAGTATGTAAACTCTGGGCTCTGTGTGTGCTGAG 98

DB 74932 GGAGTATGTAAACTCTGGGCTCTGTGTGTGCTGAG 74894

RESULT 15

AC034249

LOCUS

DEFINITION

AC034249

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

AC034249

|||||
Db 3599 GAGTATGTAAACTCTGGGTCTCTGTGTGTGCTGAGT 3637

Search completed: April 25, 2003, 00:41:44
Job time : 1748.12 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:19:38 ; Search time 181.988 Seconds

(without alignments)
4900.271 Million cell updates/sec

Title: US-09-513-999C-3792_COPY_51_446

Perfect score: 396

Sequence: 1 atgggtgcatctttgcctt.....gamctgatatctcattga 396

Scoring table:

OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 2185239 seqs, 1125999159 residues

Word size : 0

Total number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

N.Geneseq_101002:*

1: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1980.DAT:*
2: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1981.DAT:*
3: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1982.DAT:*
4: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1983.DAT:*
5: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1984.DAT:*
6: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1985.DAT:*
7: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1986.DAT:*
8: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1987.DAT:*
9: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1988.DAT:*
10: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1989.DAT:*
11: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1990.DAT:*
12: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1991.DAT:*
13: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1992.DAT:*
14: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1993.DAT:*
15: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1994.DAT:*
16: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1995.DAT:*
17: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1996.DAT:*
18: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1997.DAT:*
19: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1998.DAT:*
20: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1999.DAT:*
21: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2000.DAT:*
22: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2001A.DAT:*
23: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
24: /SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	396	100.0	447	21 AAC03794	Human secreted pro
2	396	100.0	447	21 AAZ42680	Human 5' EST isola
3	29	7.3	966	23 AAS93299	DNA encoding novel
4	24	6.1	660	23 AAS73441	DNA encoding novel
5	22	5.6	3659	7 AAN60204	Interferon-pseudo-
6	22	5.6	147724	24 ABR83566	Human cDNA differe
7	20	5.1	1179	23 AAS91061	DNA encoding novel
8	20	5.1	2526	23 AAS66674	DNA encoding novel
9	19	4.8	366	22 AAC90702	Human secretory pr

C	10	19	4.8	802	20	AAX84949	Human secreted pro
C	11	19	4.8	1982	21	AAC68089	Human secreted pro
C	12	19	4.8	2770	23	ABL09112	Drosophila melanog
C	13	19	4.8	4902	23	ABL28930	Drosophila melanog
C	14	18	4.5	31	16	AAT45064	21-hydroxylase fra
C	15	18	4.5	339	21	AAZ43080	C. elegans insulin
C	16	18	4.5	431	24	ABK24383	DNA encoding human
C	17	18	4.5	431	24	ABK24390	DNA encoding human
C	18	18	4.5	432	22	AAI83964	Human polynucleoti
C	19	18	4.5	570	22	ABA63453	Human foetal liver
C	20	18	4.5	570	22	ABA30652	Probe #9118 for ge
C	21	18	4.5	570	22	AAK11985	Human brain expres
C	22	18	4.5	570	22	AAK37688	Human bone marrow
C	23	18	4.5	570	22	AAI18447	Probe #8380 for ge
C	24	18	4.5	570	22	AAI43563	Probe #12249 used
C	25	18	4.5	570	24	ABS11680	Human genome-deriv
C	26	18	4.5	827	23	AAS87115	DNA encoding novel
C	27	18	4.5	852	22	AAI94104	Human neuroblastom
C	28	18	4.5	870	22	AAI94266	Human neuroblastom
C	29	18	4.5	872	21	AACT9760	Human secreted pro
C	30	18	4.5	3342	23	AAS87118	DNA encoding novel
C	31	18	4.5	4886	22	AAK85823	Human immune/haema
C	32	18	4.5	6464	22	ABA09665	Human bone marrow
C	33	18	4.5	9899	22	AAK85825	Human immune/haema
C	34	18	4.5	10918	22	ABA09581	Human bone marrow
C	35	18	4.5	10920	22	ABN60009	Novel human coding
C	36	18	4.5	113515	24	ABL34174	Human immune syste
C	37	17	4.3	266	20	AAV89709	EST clone CT857.
C	38	17	4.3	288	21	AACT70647	Single nucleotide
C	39	17	4.3	288	21	AACT70650	Single nucleotide
C	40	17	4.3	288	21	AACT70656	Single nucleotide
C	41	17	4.3	288	21	AACT70658	Single nucleotide
C	42	17	4.3	288	21	AACT70671	Single nucleotide
C	43	17	4.3	294	22	AAH73108	Human cervical can
C	44	17	4.3	297	22	AAH70408	Human cervical can
C	45	17	4.3	318	22	AAH69668	Human cervical can

ALIGNMENTS

RESULT 1	
AAC03794	standard; cDNA: 447 BP.
ID	AAC03794
XX	
AC	AAC03794;
XX	
DT	06-OCT-2000 (first entry)
XX	
DE	Human secreted protein 5' EST, SEQ ID NO: 3792.
XX	
XX	Human; 5' EST: expressed sequence tag; secreted protein; cDNA isolation;
KW	gene therapy; chromosome mapping; ss.
KW	
XX	
OS	Homo sapiens.
XX	
PN	EP103401-A2.
XX	
PD	06-SEP-2000.
XX	
PF	21-FEB-2000; 2000EP-0200610.
XX	
PR	26-FEB-1999; 99US-0122487.
XX	
PA	(GEST) GENSET.
PI	Dunas Milne Edwards J, Duclert A, Giordano J;
XX	
DR	WPI; 2000-500381/45.
XX	
PT	P-PSDB; AAG03788.
XX	
PT	New nucleic acid that is a 5' expressed sequence tag (5' EST) for obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for

RESULT 3
ID AAS93299 standard; cDNA: 966 BP.
XX AAS93299;
AC AAS93299;
XX 13-FEB-2002 (first entry)
XX 13-FEB-2002 (first entry)
XX DNA encoding novel human diagnostic protein #29103.
XX Human: chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX Homo sapiens.
XX WO200175067-A2.
XX 11-OCT-2001.
XX 30-MAR-2001; 2001WO-US08631.
XX 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
XX (HYSE-) HYSEQ INC.
XX Drmanac RT, Liu C, Tang YT;
XX WPI: 2001-639362/73.
XX P-PSDB: ABG29112.
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity -
XX Claim 1; SEQ ID No 29103; 103pp; English.
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. AAS64197-AAS94564 represent novel human
XX diagnostic coding sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences.
XX Sequence 966 BP; 250 A; 221 C; 259 G; 236 T; 0 other;
XX
XX Query Match 7.3%; Score 29; DB 23; Length 966;
XX Best Local Similarity 100.0%; Pred. No. 0.00027;
XX Matches 29; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX 197 CAAAGATCTGTGGAGAAAGTGTGTTCC 225
XX |||||||||||||||||||
XX 211 CAAAGATCTGTGGAGAAAGTGTGTTCC 239

RESULT 4
ID AAS73441/C
XX AAS73441 standard; cDNA: 660 BP.
XX AAS73441;
AC AAS73441;
XX 13-FEB-2002 (first entry)
XX 13-FEB-2002 (first entry)
XX DNA encoding novel human diagnostic protein #9245.
XX Human: chromosome mapping; gene mapping; gene therapy; forensic;
XX food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX Homo sapiens.
XX WO200175067-A2.
XX 11-OCT-2001.
XX 30-MAR-2001; 2001WO-US08631.
XX 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
XX (HYSE-) HYSEQ INC.
XX Drmanac RT, Liu C, Tang YT;
XX WPI: 2001-639362/73.
XX P-PSDB: ABG09254.
XX New isolated polynucleotide and encoded polypeptides, useful in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits and to assess
XX biodiversity -
XX Claim 1; SEQ ID No 9245; 103pp; English.
XX The invention relates to isolated polynucleotide (I) and
XX polypeptide (II) sequences. (I) is useful as hybridisation probes,
XX polymerase chain reaction (PCR) primers, oligomers, and for chromosome
XX and gene mapping, and in recombinant production of (II). The
XX polynucleotides are also used in diagnostics as expressed sequence tags
XX for identifying expressed genes. (I) is useful in gene therapy techniques
XX to restore normal activity of (II) or to treat disease states involving
XX (II). (II) is useful for generating antibodies against it, detecting or
XX quantitating a polypeptide in tissue, as molecular weight markers and as
XX a food supplement. (II) and its binding partners are useful in medical
XX imaging of sites expressing (II). (I) and (II) are useful for treating
XX disorders involving aberrant protein expression or biological activity.
XX The polypeptide and polynucleotide sequences have applications in
XX diagnostics, forensics, gene mapping, identification of mutations
XX responsible for genetic disorders or other traits to assess biodiversity
XX and to produce other types of data and products dependent on DNA and
XX amino acid sequences. AAS64197-AAS94564 represent novel human
XX diagnostic coding sequences of the invention.
XX Note: The sequence data for this patent did not appear in the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences.
XX Sequence 660 BP; 149 A; 127 C; 148 G; 236 T; 0 other;
XX
XX Query Match 6.1%; Score 24; DB 23; Length 660;
XX Best Local Similarity 100.0%; Pred. No. 0.076;
XX Matches 24; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX 69 AAACCTCTGGGTCTGTGTGTG 92
XX |||||||||||||||||||
XX 656 AAACCTCTGGGTCTGTGTGTG 633
RESULT 5
AAM60204

ID AAN60204 standard; DNA; 3659 BP.
XX
AC AAN60204;
XX
DT 05-AUG-1991 (first entry)
XX
DE Interferon-pseudo-omega-4.
XX
KW Interferon-omega; virucide; antitumor; ss.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT CDS 2951..3251
FT /tag= a
FT /label= interferon-pseudo-omega-4
XX
EN EPI70204-A.
XX
XX 05-FEB-1986.
XX
XX 24-JUL-1985; 85EP-0109284.
XX
XX 14-FEB-1985; 85DE-3505060.
XX 01-AUG-1984; 84DE-3428370.
XX
XX (BOEH) BOEHRINGER INGELHEIM.
XX
XX Hauptmann R, Meindl P, Dworkin-Rastl E, Adolf G, Sweetly P;
XX Pieler C, Hanel N;
XX WPI: 1986-036962/06.
XX DR P-PSDB: AAP60256.
XX
XX New interferon omega polypeptide derivs. - useful as antiviral
XX PT and antitumor agents, and new DNA sequences and genes coding for
XX PT them.
XX
XX PS Disclosure; Fig 14; 115pp; German.
XX
XX CC The sequence encodes an interferon analogue, which has virucide
XX CC and antitumor activity, and shows a synergistic increase in
XX CC activity with interferon-gamma.
XX
XX SQ Sequence 3659 BP; 796 A; 885 C; 1039 G; 939 T; 0 other;
XX
XX Query Match 5.6%; Score 22; DB 7; Length 3659;
XX Best Local Similarity 100.0%; Pred. No. 0.7;
XX Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX OY 194 TTGCAAGATCTGTGAGAG 215
XX ||||||||||||||||||||
XX Db 2557 TTGCAAGATCTGTGAGAG 2578
XX
XX RESULT 6
XX ABR83566/c
XX ID ABR83566 standard; cDNA; 147724 BP.
XX
XX AC ABR83566;
XX
XX 14-AUG-2002 (first entry)
XX
XX DE Human cDNA differentially expressed in granulocytic cells #137.
XX
XX XX Human; ss; granulocytic cell; DNA chip; bacterial infection;
XX KM viral infection; parasitic infection; protozoal infection;
XX KM fungal infection; sterile inflammatory disease; psoriasis;
XX KM rheumatoid arthritis; glomerulonephritis; asthma; thrombosis;
XX KM cardiac reperfusion injury; renal reperfusion injury; ARDS;
XX KM adult respiratory distress syndrome; inflammatory bowel disease;
XX KM Crohn's disease; ulcerative colitis; periodontal disease;
XX KM granulocyte activation; chronic inflammation; allergy.

XX
XX OS Homo sapiens.
XX
XX MO200228999-A2.
XX
XX PD 11-APR-2002.
XX
XX PF 03-OCT-2001; 2001WO-US30821.
XX
XX PR 03-OCT-2000; 2000US-237189P.
XX
XX (GENE-) GENE LOGIC INC.
XX
XX PA Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;
XX
XX DR WPI: 2002-435328/46.
XX
XX PT Detecting granulocyte activation by detecting differential expression
XX PT of genes associated with granulocyte activation, which serves as
XX PT diagnostic markers that is useful for monitoring disease states and
XX PT drug toxicity
XX
XX Claim 1; SEQ ID NO 137; 114pp; English.
XX
XX CC The invention relates to detecting (M1) granulocyte (GC) activation
XX CC (GCA), by detecting the level of expression of gene(s) (Gs) identified by
XX CC DNA chip analysis as given in the specification, and comparing
XX CC the expression level to an expression level in an unactivated
XX CC GC, where differential expression of Gs is indicative of GCA.
XX CC Also included are modulating (M2) GA by contacting GC with an agent
XX CC that alters the expression of at least one gene in Gs; (2) screening (M3)
XX CC for an agent capable of modulating GCA or an inflammation (especially
XX CC chronic) in a tissue, an allergic response in a subject, exposure of a
XX CC subject to a pathogen or sterile inflammatory disease using the
XX CC gene expression profile; (3) detecting (M4) an inflammation (especially
XX CC chronic) in a tissue, an allergic response in a subject, exposure of a
XX CC subject to a pathogen or sterile inflammatory disease, by detecting the
XX CC level of expression in a sample of the tissue of gene(s) from Gs, where
XX CC the level of expression of the gene is indicative of inflammation;
XX CC (4) treating (M5) an inflammation (especially chronic) or in a tissue,
XX CC an allergic response in a subject, exposure of a subject to a pathogen
XX CC or sterile inflammatory disease, by contacting a tissue having
XX CC inflammation with an agent that modulates the expression of gene(s)
XX CC from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for
XX CC modulating GCA; M3 is useful for screening an agent capable of modulating
XX CC GCA preferably in an inflammation in a tissue; M4 is useful for
XX CC detecting an inflammation (especially chronic) in a tissue, an allergic
XX CC response in a subject, exposure of a subject to a pathogen or sterile
XX CC inflammatory disease (e.g. psoriasis, rheumatoid arthritis,
XX CC glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal
XX CC reperfusion injury, ARDS, adult respiratory distress syndrome,
XX CC inflammatory bowel disease, Crohn's disease, ulcerative colitis,
XX CC periodontal disease; also bacterial infection, viral infection,
XX CC parasitic infection, protozoal infection, fungal infection and M5 is
XX CC useful for treating one of the above conditions. The present
XX CC sequence represents a gene differentially expressed in granulocytes.
XX CC Note: The sequence data for this patent did not form part
XX CC of the printed specification, but was obtained in electronic
XX CC format directly from WIPO at
XX CC ftp.wipo.int/pub/published_pcl_sequences.
XX
XX SQ Sequence 147724 BP; 46968 A; 29251 C; 28325 G; 43180 T; 0 other;
XX
XX Query Match 5.6%; Score 22; DB 24; Length 147724;
XX Best Local Similarity 100.0%; Pred. No. 0.66;
XX Matches 22; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX OY 149 GGCCCTGTGTCATGGCTCAC 170
XX ||||||||||||||||||||
XX Db 123545 GGCCCTGTGTCATGGCTCAC 123524
XX
XX RESULT 7

AA91061/c
ID AA91061 standard; cDNA: 1179 BP.
XX
AC AA91061;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #2685.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KM food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
XX (HYSE-) HYSEQ INC.
XX
PI Dmanac RT, Liu C, Tang YT;
XX
PI WPI: 2001-639362/73.
XX P-PSDB: ABC26874.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX
PS Claim 1; SEQ ID No 2685; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AA64197-AA94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 1179 BP; 348 A; 292 C; 315 G; 224 T; 0 other;

Query Match 5.1%; Score 20; DB 23; Length 1179;
Best Local Similarity 100.0%; Pred. No. 6.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 191 GGGTGCAGAGATCTGTGGG 210
|||||
DB 284 GGGTGCAGAGATCTGTGGG 265

RESULT 8
AA66674
ID AA66674 standard; cDNA: 2526 BP.

XX
AC AA66674;
XX
DT 13-FEB-2002 (first entry)
XX
DE DNA encoding novel human diagnostic protein #2478.
XX
KW Human; chromosome mapping; gene mapping; gene therapy; forensic;
KM food supplement; medical imaging; diagnostic; genetic disorder; ss.
XX
OS Homo sapiens.
XX
PN WO200175067-A2.
XX
PD 11-OCT-2001.
XX
PF 30-MAR-2001; 2001WO-US08631.
XX
PR 31-MAR-2000; 2000US-0540217.
XX 23-AUG-2000; 2000US-0649167.
XX (HYSE-) HYSEQ INC.
XX
PI Dmanac RT, Liu C, Tang YT;
XX
PI WPI: 2001-639362/73.
XX P-PSDB: ABC02487.
XX
PT New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity -
XX
PS Claim 1; SEQ ID No 2478; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AA64197-AA94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
SQ Sequence 2526 BP; 871 A; 536 C; 541 G; 578 T; 0 other;

Query Match 5.1%; Score 20; DB 23; Length 2526;
Best Local Similarity 100.0%; Pred. No. 6.6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 32 TTTCATCTTTCAGGAGACTT 51
|||||
DB 2377 TTTCATCTTTCAGGAGACTT 2396

RESULT 9
AAC90702/c
ID AAC90702 standard; cDNA: 366 BP.
XX
AC AAC90702;

```

XX 14-MAR-2001 (first entry)
DT
XX
DE Human secretory protein TSC-456 nucleotide sequence SEQ ID NO:17.
XX
KW Human; secretory protein; cancer; immune disease; infectious disease;
KW lung function disorder; liver function disorder; antiinflammatory;
KW gastrointestinal disorder; cytostatic; haematopoietic; anticoagulant;
KW immunomodulatory; hepatotropic; cell proliferation-stimulant;
KW cell migratory agent; cell differentiation-inducer; ss.
XX
OS Homo sapiens.
XX
PN WO200071581-A1.
XX
PD 30-NOV-2000.
XX
PE 19-MAY-2000; 2000WO-JP03221.
XX
XX 20-MAY-1999; 99JP-0140229.
XX
XX (TAKE ) TAKEDA CHEM IND LTD.
XX
XX Itoh Y, Mogi S, Tanaka H, Ohkubo S, Ogi K,
XX WPI: 2001-032023/04.
XX DR P-PSDB: AAB36662.
XX
XX Novel secretory protein and its salt with e.g. anti-cancer,
XX anti-inflammatory and hematopoietic, effects, applicable as drugs in
XX remedies and preventives to treat diseases like cancer and immune
XX diseases
XX
XX Example 2; Page 97-98; 122pp; Japanese.
XX
XX AAC90701 to AAC90715 encode the human secretory proteins given in
XX AAB36661 to AAB36675. The proteins can have cytostatic,
XX anti-inflammatory, haematopoietic, anti-coagulant, immunomodulatory and
XX hepatotropic activities, and can be used as cell migratory agents, cell
XX proliferation-stimulants and cell differentiation-inducers. The proteins
XX are useful in the treatment and prevention of diseases such as cancer,
XX lung function disorder, liver function disorder, gastrointestinal
XX disorder and immune diseases. AAC90716 to AAC90755 represent PCR primers
XX which are used in the exemplification of the present invention.
XX
XX Sequence 366 BP; 99 A; 97 C; 83 G; 87 T; 0 other;
XX
XX Query Match 4.8%; Score 19; DB 22; Length 366;
XX Best Local Similarity 100.0%; Pred. No. 21;
XX Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 217 GTGCTTCTCGATGGGCT 235
XX |
XX 238 GTGCTTCTCGATGGGCT 220
XX
XX
XX RESULT 10
XX AAX84949/C
XX ID AAX84949 standard; DNA; 802 BP.
XX
XX AAX84949;
XX
XX 30-JUL-1999 (first entry)
XX
XX Human secreted protein gene No. 17.
XX
XX Human; secreted protein; fusion protein; gene therapy; protein therapy;
XX diagnosis; tissue; cancer; tumour; neurodegenerative disorder; leukaemia;
XX developmental abnormality; foetal deficiency; blood; allergy; renal; ds;
XX immune system; asthma; lymphocytic disease; brain; hepatic; lymphoma;
XX inflammation; ischaemic shock; Alzheimer's disease; restenosis; AIDS;
XX cognitive disorder; schizophrenia; prostate; obesity; osteoclast; thymus;
XX osteoporosis; arthritis; testis; lung; thyroiditis; thyroid; digestion;

```

```

KW endocrine; metabolism; regulation; malabsorption; gastritis; neoplasm.
XX
XX Homo sapiens.
XX
XX WO9924836-A1.
XX
XX 20-MAY-1999.
XX
XX 04-NOV-1998; 98WO-US23435.
XX
XX 17-NOV-1997; 97US-0066100.
XX PR 07-NOV-1997; 97US-0064900.
XX PR 07-NOV-1997; 97US-0064908.
XX PR 07-NOV-1997; 97US-0064911.
XX PR 07-NOV-1997; 97US-0064912.
XX PR 07-NOV-1997; 97US-0064983.
XX PR 07-NOV-1997; 97US-0064984.
XX PR 07-NOV-1997; 97US-0064985.
XX PR 07-NOV-1997; 97US-0064987.
XX PR 07-NOV-1997; 97US-0066090.
XX PR 17-NOV-1997; 97US-0066094.
XX PR 17-NOV-1997; 97US-0066095.
XX PR 17-NOV-1997; 97US-0066089.
XX
XX (HUMA-) HUMAN GENOME SCI INC.
XX
XX Carter KC, Ebner R, Endress GA, Feng P, Janat F;
XX PI Ryan H, Lafileur DW, Moore PA, Ni J, Olsen HS, Rosen CA;
XX PI Ruben SM, Shi Y, Soppet DR, Wei Y;
XX DR WPI: 1999-337740/28.
XX DR P-PSDB: AAY27583.
XX
XX New human secreted proteins and coding sequences useful for treating
XX disorders of the immune system and hyperproliferative disorders
XX
XX Claim 1; Page 290; 507pp; English.
XX
XX This sequence represents a nucleic acid molecule which encodes a
XX secreted human protein. The gene number is given in the descriptor line.
XX The gene can be used to generate fusion proteins by linking to the gene
XX to a human immunoglobulin Fc portion (e.g. AAX84924) for increasing the
XX stability of the fused protein as compared to the human protein only.
XX The invention relates to 125 novel genes and their fragments (nucleic
XX acid sequences: AAX84933-X85057; amino acid sequences AAY27567-Y27933)
XX which are useful for preventing, treating or ameliorating medical
XX conditions e.g. by protein or gene therapy. Also, pathological
XX conditions can be diagnosed by determining the amount of the new
XX polypeptides in a sample or by determining the presence of mutations in
XX the new polynucleotides. Specific uses are described for each of the 125
XX polynucleotides, based on which tissues they are most highly expressed in
XX (see AAX84933 for described uses).
XX
XX Sequence 802 BP; 208 A; 210 C; 190 G; 188 T; 6 other;
XX
XX Query Match 4.8%; Score 19; DB 20; Length 802;
XX Best Local Similarity 100.0%; Pred. No. 21;
XX Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
XX
XX QY 217 GTGCTTCTCGATGGGCT 235
XX |
XX Db 304 GTGCTTCTCGATGGGCT 286
XX
XX
XX RESULT 11
XX AAC68089/C
XX ID AAC68089 standard; CDNA; 1982 BP.
XX
XX AAC68089;
XX
XX 20-FEB-2001 (first entry)
XX

```


DE Human secreted protein cDNA sequence #9.
 XX
 XX Cytostatic; immunosuppressive; nootropic; neuroprotective; antiviral;
 KW antiallergic; hepatotropic; antidiabetic; antiinflammatory; antitumor;
 KW vulnery; anticonvulsant; antibacterial; antifungal; antiparasitic;
 KW cardiant; gene therapy; cancer; immune disorder; cardiovascular disorder;
 KW neurological disease; infection; human; secreted protein; ss.
 XX
 XX Homo sapiens.
 OS
 XX
 XX MO200058335-A1.
 PN
 XX
 XX 05-OCR-2000.
 PD
 XX
 XX 22-MAR-2000; 2000WO-US07534.
 PE
 XX
 XX 26-MAR-1999; 99US-0126598.
 PR
 XX 22-DEC-1999; 99US-0171504.
 PR
 XX
 XX (HUMA-) HUMAN GENOME SCI INC.
 BA (ROSE/) ROSEN C A.
 XX
 XX Rosen CA, Ruben SM, Komatsoulis G;
 XI
 XX WPI: 2000-611702/58.
 DR
 XX P-PSDB; AAB37356.
 DR
 XX
 XX Nucleic acids encoding human secreted proteins, used to treat, prevent,
 PT ameliorate or diagnose conditions such as cancer, and autoimmune
 PT diseases e.g. arthritis -
 PT
 XX
 XX
 XX Claim 1; Pages 321-322; 387pp; English.
 XX
 XX The invention relates to the isolation of genes AAC68081-C68127 encoding
 CC 47 human secreted proteins AAB37348-B37394. The genes can be used to
 CC generate fusion proteins by linking to the gene for the human
 CC immunoglobulin G Fc portion (AAC68072) for increasing the stability of
 CC the fusion protein as compared to the human protein only. The genes and
 CC proteins are useful for preventing, ameliorating or treating medical
 CC conditions, e.g. by protein or gene therapy. The genes are isolated
 CC from a range of human tissues disclosed in the specification. The
 CC nucleic acids, proteins, antibodies and (ant)agonists are useful in
 CC the diagnosis, treatment and prevention of: (a) cancer, e.g. breast
 CC and ovarian cancer, and other cancers of the adrenal gland, bone, bone
 CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;
 CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune
 CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
 CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative
 CC colitis; (c) cardiovascular disorders such as myocardial ischaemias; (d)
 CC wound healing; (e) neurological diseases e.g. cerebral anoxia and
 CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal
 CC and parasitic infections.
 CC
 XX
 XX Sequence 1982 BP; 549 A; 513 C; 471 G; 448 T; 1 other;
 SQ
 XX
 XX Query Match 4.8%; Score 19; DB 21; Length 1982;
 XX Best Local Similarity 100.0%; Pred. No. 21;
 XX Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 326 TTTTCTCATCTCTGTGG 344
 XX ||||||||||||||||
 DB 860 TTTTCTCATCTCTGTGG 842
 XX
 XX
 XX RESULT 12
 XX ABL09112
 XX ID ABL09112 standard; cDNA: 2770 BP.
 XX
 XX ABL09112;
 AC
 XX 26-MAR-2002 (first entry)
 DT
 XX
 XX Drosophila melanogaster expressed polynucleotide SEQ ID NO 21818.

XX
 XX Drosophila; developmental biology; cell signalling; insecticide;
 KW pharmaceutical; gene; ss.
 XX
 XX Drosophila melanogaster.
 OS
 XX
 XX WO200171042-A2.
 PN
 XX
 XX 27-SEP-2001.
 PD
 XX
 XX 23-MAR-2001; 2001WO-US09231.
 PE
 XX
 XX 23-MAR-2000; 2000US-191637P.
 PR
 XX 11-JUL-2000; 2000US-0614150.
 PR
 XX
 XX (PEKE) PE CORP NY.
 PA
 XX
 XX Venter JC, Adams M, Li PWD, Myers EW;
 PI
 XX
 XX WPI: 2001-656860/75.
 DR
 XX P-PSDB; ABB65009.
 DR
 XX
 XX New isolated nucleic acid detection reagent for detecting 1000 or more
 PT genes from Drosophila and for elucidating cell signalling and cell-cell
 PT interactions -
 PT
 XX
 XX
 XX Claim 1; SEQ ID NO 21818; 21pp + Sequence Listing; English.
 XX
 XX The invention relates to an isolated nucleic acid detection reagent
 CC capable of detecting 1000 or more genes from Drosophila. The invention is
 CC useful in developmental biology and in elucidating cell signalling and
 CC cell-cell interactions in higher eukaryotes for the development of
 CC insecticides, therapeutics and pharmaceutical drugs. The invention
 CC discloses genomic DNA sequences (ABL16176-ABL30511), expressed DNA
 CC sequences (ABL01840-ABL16175) and the encoded proteins
 CC (ABB57737-ABB72072).
 CC
 CC The sequence data for this patent did not form part of the printed
 CC specification and was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 CC
 XX
 XX Sequence 2770 BP; 786 A; 625 C; 629 G; 730 T; 0 other;
 SQ
 XX
 XX Query Match 4.8%; Score 19; DB 23; Length 2770;
 XX Best Local Similarity 100.0%; Pred. No. 20;
 XX Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 233 GGTGACACATCTACTACT 251
 XX ||||||||||||||||
 DB 1050 GGTGACACATCTACTACT 1068
 XX
 XX
 XX RESULT 13
 XX ABL28930/c
 XX ID ABL28930 standard; DNA; 4902 BP.
 XX
 XX ABL28930;
 AC
 XX 26-MAR-2002 (first entry)
 DT
 XX
 XX Drosophila melanogaster genomic polynucleotide SEQ ID NO 38263.
 DE
 XX
 XX Drosophila; developmental biology; cell signalling; insecticide;
 KW pharmaceutical; gene; ds.
 KW
 XX
 XX Drosophila melanogaster.
 OS
 XX
 XX WO200171042-A2.
 PN
 XX
 XX 27-SEP-2001.
 PD
 XX
 XX 23-MAR-2001; 2001WO-US09231.
 PE
 XX
 XX 23-MAR-2000; 2000US-191637P.
 PR
 XX

PR 11-JUL-2000; 2000US-0614150.
 XX
 XX (PEKE) PE CORP NY.
 XX
 PI Venter JC, Adams M, Li PWD, Myers EW.
 XX
 DR WPI: 2001-656860/75.
 XX
 PT New isolated nucleic acid detection reagent for detecting 1000 or more
 PT genes from Drosophila and for elucidating cell signalling and cell-cell
 PT interactions -
 XX
 PS Claim 1; SEQ ID NO 38263; 21pp + Sequence Listing; English.
 XX
 CC The invention relates to an isolated nucleic acid detection reagent
 CC capable of detecting 1000 or more genes from Drosophila. The invention is
 CC useful in developmental biology and in elucidating cell signalling and
 CC cell-cell interactions in higher eukaryotes for the development of
 CC insecticides, therapeutics and pharmaceutical drugs. The invention
 CC discloses genomic DNA sequences (AB116176-AB130511), expressed DNA
 CC sequences (AB101840-AB116175) and the encoded proteins
 CC (ABB57737-ABB72072).
 CC The sequence data for this patent did not form part of the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 CC
 XX Sequence 4902 BP; 1509 A; 1006 C; 984 G; 1403 T; 0 other;
 SQ
 Query Match 4.8%; Score 19; DB 23; Length 4902;
 Best Local Similarity 100.0%; Pred. No. 20;
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 268 GGTGGGCTTTTGTTCG 286
 |||||||
 Db 886 GGTGGGCTTTTGTTCG 868
 RESULT 14
 AAT45064
 ID AAT45064 standard; DNA; 31 BP.
 XX
 AC AAT45064;
 XX
 DT 05-FEB-1997 (first entry)
 XX
 DE 21-hydroxylase fragment A, forward primer.
 XX
 Polymerase chain reaction; primer; PCR; amplify; 21-hydroxylase gene;
 human adrenal gland cDNA library; diagnosis; detection;
 Addison's disease; ss.
 XX
 OS Synthetic.
 XX
 PN US5376533-A.
 XX
 PD 27-DEC-1994.
 XX
 PF 24-MAY-1993; 93US-0066281.
 XX
 PR 24-MAY-1993; 93US-0066281.
 XX
 PA (UYFL) UNIV FLORIDA.
 XX
 PI MacIaren NK, Song YH;
 XX
 DR WPI: 1995-043462/06.
 XX
 PT Detection of Addison's disease or persons at risk from developing it
 PT using a 21-hydroxylase peptide fragment to detect autoantibodies
 PT associated with Addison's disease.
 XX
 PS Disclosure; Column 6; 9pp; English.
 XX

CC The sequences given in AAT45064-75 are primers which were used to
 CC amplify fragments of the 21-hydroxylase gene from a human adrenal
 CC gland cDNA library. The amplified fragments encode peptides which
 CC were used in the method of the invention to diagnose or detect
 CC Addison's disease. Fragment A comprises amino acids 1-162, fragment
 CC B, amino acids 164-356, fragment C, amino acids 3440494, fragment D,
 CC amino acids 164-271, fragment E, amino acids 272-356, and fragment F
 CC amino acids 197-298 of the 21-hydroxylase enzyme. These primers
 CC have been optimised for use and do not directly correspond to 21
 CC hydroxylase gene sequences. All the forward primers contain BamHI
 CC sites and all the reverse primers contain EcoRI restriction sites.
 XX
 SQ Sequence 31 BP; 2 A; 10 C; 11 G; 8 T; 0 other;
 SQ
 Query Match 4.5%; Score 18; DB 16; Length 31;
 Best Local Similarity 100.0%; Pred. No. 67;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 OY 288 TCCATCCTGCTCCTGGGC 305
 |||||||
 Db 8 TCCATCCTGCTCCTGGGC 25
 RESULT 15
 AA243080/C
 ID AA243080 standard; DNA; 339 BP.
 XX
 AC AA243080;
 XX
 DT 04-FEB-2000 (first entry)
 XX
 DE C. elegans insulin-like protein ZK84.6 gene.
 XX
 KW Insulin-like protein; diagnosis; insulin-like gene analysis; nematode;
 KW insulin hormone; aging; senescence; pesticide target; signalling pathway;
 KW signal transduction pathway; ss.
 XX
 OS Caenorhabditis elegans.
 XX
 PN W09954436-A2.
 XX
 PD 28-OCT-1999.
 XX
 PF 16-APR-1999; 99WO-US08522.
 XX
 PR 17-APR-1998; 98US-0062580.
 PR 08-MAY-1998; 98US-0074984.
 PR 26-MAY-1998; 98US-0084303.
 XX
 PA (EXEL-) EXELIXIS PHARM INC.
 XX
 PI Homburger SA, Platt DM, Ferguson KC, Doberstein SK, Buchman AR;
 PI Reddy BP;
 XX
 DR WPI: 2000-012239/01.
 XX
 DR P-PSDB; AAY65657.
 XX
 PT Analysing Caenorhabditis elegans insulin-like gene expression, nucleic
 PT acids and proteins of the C. elegans insulin-like genes -
 XX
 PS Claim 4; Fig 8; 194pp; English.
 XX
 CC This sequence encodes a Caenorhabditis elegans insulin-like protein,
 CC and can be used in the method of the invention. The method is for
 CC analysing an effect of expression or mis-expression of a C. elegans
 CC insulin-like gene, and comprises observing a first nematode genetically
 CC engineered to express or mis-express a C. elegans insulin-like protein
 CC (ILP) of any one of groups I, II or IV or a derivative or fragment that
 CC displays one or more functional activities of the C. elegans ILP. The
 CC insulin-like genes in C. elegans constitute very useful tools for probing
 CC the function and regulation of their corresponding pathways. This can be
 CC expected to lead to the discovery of new drug targets, therapeutic
 CC proteins, diagnostics and prognostics useful in the treatment of diseases

CC and clinical problems associated with the function of insulin hormones in
 CC humans and other animals, as well as clinical problems associated with
 CC aging and senescence. The information may also be useful in
 CC identification and validation of pesticide targets in invertebrate pests
 CC that are components of these signalling pathways. The genes are also
 CC useful for identifying factors that are upstream of the receptor in the
 CC signal transduction pathway. The ligand-encoding C. elegans insulin-like
 CC genes provide a superior approach for identifying factors that are
 CC upstream of the receptor in the signal transduction pathway.

CC
 XX Sequence 339 BP; 92 A; 86 C; 72 G; 89 T; 0 other;

Query Match 4.5%; Score 18; DB 21; Length 339;

Best Local Similarity 100.0%; Pred. No. 65;

Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 188 TGTGGTTGCCAAGATCT 205

|||||

Db 260 TGTGGTTGCCAAGATCT 243

Search completed: April 25, 2003, 00:01:03
 Job time : 220.988 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:49:04 ; Search time 1331.72 Seconds
(without alignments)
4815.901 Million cell updates/sec

Title: US-09-513-999C-3792_COPY_51_446

Perfect score: 396

Sequence: 1 atgggtgattcatttcgcctt.....gamctgatatcttcagtta 396

Scoring table: OLIGO_NUC

Gapop 60.0 , Gapext 60.0

Searched: 16154066 seqs, 8097743376 residues

d size : 0

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database :

EST:*
1: em_estba:*
2: em_esthum:*
3: em_estin:*
4: em_estmu:*
5: em_estov:*
6: em_estpl:*
7: em_estro:*
8: em_hlc:*
9: gb_est1:*
10: gb_est2:*
11: gb_hlc:*
12: gb_est3:*
13: gb_est4:*
14: gb_est5:*
15: em_estfun:*
16: em_estom:*
17: gb_gss:*
18: em_gss_hum:*
19: em_gss_inv:*
20: em_gss_pln:*
21: em_gss_vrt:*
22: em_gss_fun:*
23: em_gss_mam:*
24: em_gss_mus:*
25: em_gss_other:*
26: em_gss_pro:*
27: em_gss_rpd:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	147	37.1	470	17	A0770688 HS_5368_B
2	146	36.9	525	17	A0165256 HS_3025_B
3	38	9.6	628	17	A0237815 RPCI11-70
4	35	8.8	410	17	A0442274 HS_5137_A
5	35	8.8	453	17	A0437684 HS_5137_A
6	35	8.8	635	17	A0390599 CITBI-EL-

Result No.	Score	Query Match	Length	DB ID	Description
7	34	8.6	412	17	A0321916 RPCI11-10
8	34	8.6	529	17	A0881246 HS_5137_B
9	33	8.3	378	17	A0479650 RPCI-11-2
10	33	8.3	471	17	A0147593 HS_3065_B
11	33	8.3	482	17	A0320567 RPCI11-99
12	33	8.3	546	12	BF962702 PM4-NM120
13	33	8.3	563	17	A0420187 RPCI-11-1
14	33	8.3	691	17	A2516454 RPCI-11-1
15	33	8.3	723	17	A0386439 RPCI11-15
16	32	8.1	427	17	A0697116 HS_5528_A
17	31	7.8	553	17	A2521751 RPCI-11-1
18	31	7.8	695	17	AG179297 Pan trogl
19	31	7.8	769	17	A0899390 HS_5234_A
20	30	7.6	360	17	A0207172 HS_3238_B
21	30	7.6	399	17	A0115544 RPCI11-57
22	30	7.6	435	17	A0116061 RPCI11-57
23	30	7.6	452	17	A0442744 HS_5122_B
24	30	7.6	551	17	A0569689 HS_5333_B
25	29	7.3	414	17	A0003326 RPCI11-25
26	29	7.3	468	17	A0819715 HS_5513_A
27	29	7.3	615	17	AG161224 Pan trogl
28	29	7.3	653	17	AG143347 Pan trogl
29	28	7.1	363	17	A0120796 HS_3076_A
30	28	7.1	376	17	A0548294 RPCI-11-4
31	28	7.1	401	17	A0588089 CITBI-EL-
32	28	7.1	419	17	A036209 HS_5049_B
33	28	7.1	425	17	A0141070 HS_3141_B
34	28	7.1	449	17	A0269217 RPCI11-69
35	28	7.1	490	17	A0817757 HS_5250_B
36	28	7.1	530	17	A0193128 HS_3060_B
37	28	7.1	541	17	A0683783 HS_5455_B
38	28	7.1	653	17	AG160919 Pan trogl
39	28	7.1	681	17	AG141287 Pan trogl
40	28	7.1	684	17	AG049745 Pan trogl
41	27	6.8	345	17	A0075574 CIT-HSP-2
42	27	6.8	517	17	A0614252 HS_5123_B
43	27	6.8	529	17	A0346593 RPCI11-12
44	27	6.8	553	17	A0238365 RPCI11-63
45	27	6.8	606	17	A0350708 RPCI11-11

ALIGNMENTS

RESULT 1 A0770688 470 bp DNA linear GSS 28-JUL-1999
DEFINITION HS_5368_B2.C08.SP66 RPCI-11 Human Male BAC library Homo sapiens
LOCUS A0770688 genomic clone Plate=944 Col-16 Row=F, DNA sequence.
ACCESSION A0770688
VERSION A0770688.1 GI:5648804
KEYWORDS GSS.

ORGANISM

human.

REFERENCE

1 (bases 1 to 470)

AUTHORS

Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.

TITLE

Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

JOURNAL

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)

COMMENT

High Throughput Sequencing Center
University of Washington

CONTACT

Contact: Mahairas GG, Wallace JC, Hood L
401 Queen Anne Avenue North, Seattle, WA 98109, USA

TELEPHONE

Tel: (206) 616-3618

FAX

Fax: (206) 616-3887

EMAIL

Email: jwallace@u.washington.edu

Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong

BASE COUNT 125 a 150 c 164 g 189 t
ORIGIN
Query Match 9.6%; Score 38; DB 17; Length 628;
Best Local Similarity 100.0%; Pred. No. 2.6e-06;
Matches 38; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 61 GAGTATGTAACCTCGCTCTGTGTGCTGAG 98
|||||
Db 157 GAGTATGTAACCTCGCTCTGTGTGCTGAG 194

RESULT 4
LOCUS A0442274 410 bp DNA linear GSS 31-MAR-1999
DEFINITION HS_5137_A1_P12.SP6E RPCT-11 Human Male BAC Library Homo sapiens
ACCESSION A0442274
VERSION A0442274.1 GI:4553613
MEDLINE
JOURNAL
COMMENT human
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 410)
Mahairas G.G., Wallace J.C., Smith K., Swartzell S., Holzman T.,
Keller A., Shaker R., Furlong J., Young J., Zhao S., Adams M.D. and
Hood L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
99380589
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCT-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 713 row: K column: 23
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 410.
Location/Qualifiers
1. 410
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=713 Col=23 Row=K"
/clone_lib="RPCT-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBACe3.6 vector at EcoRI sites"

BASE COUNT 80 a 111 c 111 g 107 t 1 others
ORIGIN
Query Match 8.8%; Score 35; DB 17; Length 410;
Best Local Similarity 100.0%; Pred. No. 6e-05;
Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 TCCTGGGCTCTGTGTGTGCTGAGTGGCTGCT 108
|||||
Db 160 TCCTGGGCTCTGTGTGTGCTGAGTGGCTGCT 194

RESULT 5
LOCUS A0437684 453 bp DNA linear GSS 31-MAR-1999
DEFINITION HS_5137_A2_H06.SP6E RPCT-11 Human Male BAC Library Homo sapiens
ACCESSION A0437684
VERSION A0437684.1 GI:4549023
MEDLINE
JOURNAL
COMMENT human
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 453)
Mahairas G.G., Wallace J.C., Smith K., Swartzell S., Holzman T.,
Keller A., Shaker R., Furlong J., Young J., Zhao S., Adams M.D. and
Hood L.
Sequence-tagged connectors: A sequence approach to mapping and
scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
99380589
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPCT-11. For BAC
library availability, please contact Pieter de Jong
(pieter@dejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 713 row: O column: 12
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 453.
Location/Qualifiers
1. 453
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=713 Col=12 Row=O"
/clone_lib="RPCT-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBACe3.6 vector at EcoRI sites"

BASE COUNT 84 a 127 c 117 g 124 t 1 others
ORIGIN
Query Match 8.8%; Score 35; DB 17; Length 453;
Best Local Similarity 100.0%; Pred. No. 5.8e-05;
Matches 35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 74 TCCTGGGCTCTGTGTGTGCTGAGTGGCTGCT 108
|||||
Db 163 TCCTGGGCTCTGTGTGTGCTGAGTGGCTGCT 197

RESULT 6
LOCUS A0390599 635 bp DNA linear GSS 06-MAR-1999
DEFINITION CITBI-El-2544B15.TR CITBI-El Homo sapiens genomic clone 2544B15,
DNA sequence.
ACCESSION A0390599
VERSION A0390599.1 GI:4361622
MEDLINE
JOURNAL
COMMENT human
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE	Mammalia: Eutheria: Primates: Catarrhini: Homnidae; Homo.		
AUTHORS	1 (bases 1 to 635) Zhao,S., Adams,M.D., Niernan,W., Malek,J., Shizuya,H., Simon,M. and Venter,J.C.		
TITLE	Use of BAC End Sequences from Caltech Libraries for Sequence-Ready Map Building		
JOURNAL	Unpublished (1997)		
COMMENT	Other.GSS: CITBI-EI-2544B15.TF Contact: Shaying Zhao, William Niernan, Mark Adams Department of Eukaryotic Research The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850 Tel: 301 838 0200 Fax: 301 838 0208 Email: hbeetlgr.org Clones are available from Research Genetics (inforesgen.com). BAC end search page: http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html . Seq primer: M13 Reverse Class: BAC ends.		
URES	Location/Qualifiers		
source	1..635 /organism="Homo sapiens" /db_xref="taxon:9606" /clone="2544B15" /clone_11b="CITBI-EI" /sex="male" /cell_type="sperm" /note="Vector: pBeloBAC11; Site_1: EcoRI; Site_2: EcoRI; Caltech Human BAC Library D"		
BASE COUNT	192 a 144 c 172 g 127 t		
ORIGIN			
Query Match	8.8%; Score 35; DB 17; Length 635;		
Best Local Similarity	100.0%; Pred. No. 5,4e-05;		
Matches	35; Conservative 0; Mismatches 0; Indels 0; Gaps 0;		
QY	74 TCCCTGGCTCTGCTGTGCTGCTGAGTGGCTGCTCT 108 		
Db	527 TCCCTGGCTCTGCTGTGCTGCTGAGTGGCTGCTCT 493		
RESULT 7			
A0321916			
LOCUS	A0321916 412 bp DNA linear GSS 06-MAY-1999		
DEFINITION	RPC111-101H18.TJ RPC1-11 Homo sapiens genomic clone RPC1-11-101H18, DNA sequence.		
ACCESSION	A0321916		
VERSION	A0321916.1 GI:4054584		
WORDS	GSS.		
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.		
AUTHORS	1 (bases 1 to 412) Adams,M.D., Rounsley,S.D., Zhao,S., Baas,S., Linher,K., Golden,K., Barry,K., Granger,D., Sub,E., White,C., de Jong,P. and Venter,J.C.		
TITLE	Use of human BAC end sequences for Sequence-Ready Map Building		
JOURNAL	Unpublished (1998)		
COMMENT	Other.GSS: RPC111-101H18.TJ Contact: Shaying Zhao, William Niernan, Mark Adams Department of Eukaryotic Research The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850 Tel: 301 838 0200 Fax: 301 838 0208 Email: hbeetlgr.org Clones are derived from the human BAC library RPC1-11. For BAC library availability, please contact Plier de Jong (plierdejong.med.buffalo.edu). Clones may be purchased from BACRPC Resources (http://bacpac.med.buffalo.edu/ordering) or from Research Genetics (inforesgen.com). BAC end search page: http://www.tigr.org/tdb/hungen/bac_end_search/bac_end_search.html		

```

Seq primer: SP6
Class: BAC ends.

FEATURES
    source
        location/Qualifiers
            1..412
                /organism="Homo sapiens"
                /db_xref="GDB:7538585"
                /db_xref="taxon:9606"
                /clone="RPCI-11-101H18"
                /clone_11b="RPCI-11"
                /sex="Male"
                /cell_type="Lymphocytes"
                /note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
                RPC11 Human Male BAC library"
BASE COUNT
    74 a 102 c 120 g 116 t
ORIGIN
Query Match 8.6%; Score 34; DB 17; Length 412;
Best Local Similarity 100.0%; Pred. No. 0.00015;
Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 191 GGGTTGCAACAGATCTGTGGAGACAGTGTGCTTTC 224
|||||
Db 292 GGGTTGCAACAGATCTGTGGAGACAGTGTGCTTTC 325

RESULT 8
AC0881246/c 529 bp DNA linear GSS 09-NOV-1999
LOCUS HS_5137_B1_F08_T7 RPCI-11 Human Male BAC library Homo sapiens
DEFINITION genomic clone Plate=8905 Col=15 Row=L, DNA sequence.
ACCESSION AC0881246
VERSION AC0881246.1 GI:6312713
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 529)
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J.J., Young,J., Zhao,S., Adams,M.D. and
Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and
screening the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
99380839
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallaceu.washington.edu
Clones are derived from the human BAC library RPCI-11. For BAC
library availability, please contact Pieter de Jong
(pieterdejong.med.buffalo.edu). Clones may be purchased from
BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Resear h Genetics (info@resgen.com). BAC end Web Server:
http://www.htsc.washington.edu
Plate: 8905 row= L column= 15
Seq primer: T7
Class: BAC ends
High quality sequence stop: 529.

FEATURES
    source
        location/Qualifiers
            1..529
                /organism="Homo sapiens"
                /db_xref="taxon:9606"
                /clone="Plate=8905 Col=15 Row=L"
                /clone_11b="RPCI-11 Human Male BAC library"
                /sex="male"
                /note="Vector: pBACe3.6; Site_1: EcoRI; Site_2: EcoRI;
                Male blood DNA was isolated from one randomly chosen donor
                and partially digested with a combination of EcoRI and
                EcoRI Methylase. Size selected DNA was cloned into the

```


BASE COUNT 137 a 147 c 121 g 117 t 7 others
 ORIGIN PBACE3.6 vector at EcoRI sites"

Query Match 8.6%; Score 34; DB 17; Length 529;
 Best Local Similarity 100.0%; Pred. No. 0.00016;
 Matches 34; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 60 GGAGTATGTAACCTCGGCTCTGTGTGC 93
 DB 382 GGAGTATGTAACCTCGGCTCTGTGTGC 349

RESULT 9
 LOCUS AQ479650/c 378 bp DNA linear GSS 23-APR-1999
 DEFINITION RPCI-11-269D19.TV RPCI-11 Homo sapiens genomic clone RPCI-11-269D19
 , DNA sequence.
 ACCESSION AQ479650
 VERSION AQ479650.1 GI:4661769
 WORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 378)
 AUTHORS Zhao,S., Adams,M.D., Nierman,W., Malek,J., de Jong,P. and Venter,J.C.
 TITLE Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready
 Map Building
 JOURNAL Unpublished (1997)
 COMMENT Contact: Shaying Zhao, William Nierman, Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbeetlgr.org
 Clones are derived from the human BAC library RPCI-11. For BAC
 library availability, please contact Pieter de Jong
 (pieter@edjlong.med.buffalo.edu). Clones may be purchased from
 BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from
 Research Genet. cs (info@resgen.com). BAC end search page:
 http://www.tlgr.org/tldb/hungun/bac_end_search/bac_end_search.html.
 Seq primer: 17
 Class: BAC ends.

FEATURES
 Location/Qualifiers
 1..378
 /organism="Homo sapiens"
 /db_xref="GDB:7603002"
 /db_xref="taxon:9606"
 /clone="RPCI-11-269D19"
 /clone.lib="RPCI-11"
 /sex="Male"
 /cell_type="Lymphocytes"
 /note="Vector: PBACE3.6; Site_1: EcoRI; Site_2: EcoRI;
 RPCI11 Human Male BAC Library"

BASE COUNT 116 a 103 c 93 g 66 t

Query Match 8.3%; Score 33; DB 17; Length 378;
 Best Local Similarity 100.0%; Pred. No. 0.00046;
 Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 193 GTTCAAGATCTGTGGAGAGTGTTC 225
 DB 208 GTTCAAGATCTGTGGAGAGTGTTC 176

RESULT 10
 LOCUS AQ147593/c 471 bp DNA linear GSS 08-OCT-1998
 DEFINITION HS_3065_B2_H08_MF CIT Approved Human Genomic Sperm Library D Homo

ACCESSION sapiens genomic clone Plate=3065 Col=16 Row=P, DNA sequence.
 VERSION AQ147593
 KEYWORDS AQ147593.1 GI:3538246
 GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 471)
 AUTHORS Mahalas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
 Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and
 Hood,L.
 TITLE Sequence-tagged connectors: A sequence approach to mapping and
 scanning the human genome
 JOURNAL Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
 MEDLINE 99380589
 COMMENT Contact: Mahalas GG, Wallace JC, Hood L
 High Throughput Sequencing Center
 University of Washington
 401 Queen Anne Avenue North, Seattle, WA 98109, USA
 Tel: (206) 616-3618
 Fax: (206) 616-3887
 Email: jwallace@u.washington.edu
 Sequence Tagged Connector
 Plate: 3065 row: P Column: 16
 Class: BAC ends
 High quality sequence stop: 471.
 Location/Qualifiers
 1..471
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /clone="Plate=3065 Col=16 Row=P"
 /clone.lib="CIT Approved Human Genomic Sperm Library D"
 /sex="male"
 /note="Organ: sperm; Vector: pBelobAC11; BAC Clones in
 E-Coli DH10B"

BASE COUNT 144 a 118 c 123 g 84 t 2 others

Query Match 8.3%; Score 33; DB 17; Length 471;
 Best Local Similarity 100.0%; Pred. No. 0.00044;
 Matches 33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 193 GTTCAAGATCTGTGGAGAGTGTTC 225
 DB 265 GTTCAAGATCTGTGGAGAGTGTTC 233

RESULT 11
 LOCUS AQ320567/c 482 bp DNA linear GSS 04-MAY-1999
 DEFINITION RPCI11-99N1.TV RPCI-11 Homo sapiens genomic clone RPCI-11-99N1, DNA
 sequence.
 ACCESSION AQ320567
 VERSION AQ320567.1 GI:4050696
 WORDS GSS.
 SOURCE human.
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 482)
 AUTHORS Adams,M.D., Rounsley,S.D., Zhao,S., Baas,S., Linher,K., Golden,K.,
 Berry,K., Granger,D., Suh,E., Wible,C., de Jong,P. and Venter,J.C.
 TITLE Use of human BAC End Sequences for Sequence-Ready Map Building
 JOURNAL Unpublished (1998)
 COMMENT Contact: Shaying Zhao, William Nierman, Mark Adams
 Department of Eukaryotic Genomics
 The Institute for Genomic Research
 9712 Medical Center Dr., Rockville, MD 20850
 Tel: 301 838 0200
 Fax: 301 838 0208
 Email: hbeetlgr.org
 Clones are derived from the human BAC library RPCI-11. For BAC

Db 333 TGGGGCCGAGTATGTAACCTCTGGTCTCT 301.

RESULT 14

LOCUS

A2516454/c

DEFINITION

691 bp DNA linear GSS 16-OCT-2000
RPC1-11-191D19.TV RPC1-11 Homo sapiens genomic clone RPC1-11-191D19
, DNA sequence.

ACCESSION

A2516454

VERSION

A2516454.1 GI:10824878

KEYWORDS

GSS.

SOURCE

human.

ORGANISM

Homo sapiens

Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 691)

REFERENCE

AUTHORS

Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter
,J.C.

TITLE

BAC end sequences of library RPC1-11

JOURNAL

Unpublished (1997)

COMMENT

Other_GSSs: RPC1-11-191D19.TJB

FEATURES

Contact: Shaying Zhao

LOCATION

Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA

TELEPHONE

Tel: 301 838 0200

FAX

Fax: 301 838 0200

EMAIL

Email: szhao@tigr.org

CLONES

Clones are derived from the human BAC library RPC1-11. For BAC

LIBRARY

library availability, please contact Pieter de Jong

BAC

BAC PAC Resources (http://bacpac.med.buffalo.edu). Clones may be purchased from

RESEARCH

Research Genet cs (info@resgen.com). BAC end search page:

HTTP

http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html.

THIS

This BAC end was generated during the RAD process and may have

HIGHER

higher chance of clone tracking errors.

SEQ

Seq primer: 77

CLASS

Class: BAC ends.

FEATURES

Location/Qualifiers

SOURCE

1..691

ORGANISM

/organism="Homo sapiens"

DB

/db_xref="GDB:7573050"

/DB

/db_xref="taxon:9606"

/CLONE

/clone="RPC1-11-191D19"

/CLONE

/clone_lib="RPC1-11"

/SEX

/sex="Male"

/CELL

/cell_type="Lymphocytes"

/NOTE

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC111 Human Male BAC library"

BASE

COUNT

203 a 191 c 174 g 121 t 2 others

ORIGIN

Query Match

8.3%; Score 33; DB 17; Length 691;

Best

Best Local Similarity 100.0%; Pred. No. 0.0004;

Matches

33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

193 GTTGCAGAGATCTGTGGAGAGTGTGTTCC 225

DB

216 GTTGCAGAGATCTGTGGAGAGTGTGTTCC 184

RESULT 15

LOCUS

AQ386439/c

DEFINITION

723 bp DNA linear GSS 21-MAY-1999
RPC11-154D6.TJ RPC1-11 Homo sapiens genomic clone RPC1-11-154D6,
DNA sequence.

ACCESSION

AQ386439

VERSION

AQ386439.1 GI:4357462

KEYWORDS

GSS.

SOURCE

human.

ORGANISM

Homo sapiens

MAMMALIA

Mammalia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

EUTHERIA

Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE 1 (bases 1 to 723)

AUTHORS

Zhao,S., Adams,M.D., Niernan,W., Malek,J., de Jong,P. and Venter
,J.C.

TITLE

Use of BAC End Sequences from Library RPC1-11 for Sequence-Ready

JOURNAL

Map Building

COMMENT

Unpublished (1997)

Other

Other_GSSs: RPC11-154D6.TV

Contact

Contact: Shaying Zhao, William Niernan, Mark Adams

Department

Department of Eukaryotic Genomics

The

The Institute for Genomic Research

9712

9712 Medical Center Dr., Rockville, MD 20850

Tel:

Tel: 301 838 0200

Fax:

Fax: 301 838 0200

Email:

Email: hbeet@tigr.org

CLONES

Clones are derived from the human BAC library RPC1-11. For BAC

LIBRARY

library availability, please contact Pieter de Jong

BAC

BAC PAC Resources (http://bacpac.med.buffalo.edu). Clones may be purchased from

RESEARCH

Research Genet cs (info@resgen.com). BAC end search page:

HTTP

http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html

Seq

Seq primer: SP6

CLASS

Class: BAC ends.

FEATURES

Location/Qualifiers

SOURCE

1..723

ORGANISM

/organism="Homo sapiens"

DB

/db_xref="GDB:7558829"

/DB

/db_xref="taxon:9606"

/CLONE

/clone="RPC1-11-154D6"

/CLONE

/clone_lib="RPC1-11"

/SEX

/sex="Male"

/CELL

/cell_type="Lymphocytes"

/NOTE

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;
RPC111 Human Male BAC library"

BASE

COUNT

224 a 165 c 135 g 199 t

ORIGIN

Query Match

8.3%; Score 33; DB 17; Length 723;

Best

Best Local Similarity 100.0%; Pred. No. 0.0004;

Matches

33; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY

53 TGGGGCCGAGTATGTAACCTCTGGGCTCT 85

DB

331 TGGGGCCGAGTATGTAACCTCTGGGCTCT 299

Search completed: April 25, 2003, 00:52:51

Job time : 1334.72 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 24, 2003, 23:57:29 ; Search time 46.8639 Seconds

(without alignments)
2591.419 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_446

Sequence: 1 atgggtgacatcttcgcctt.....gamctgatacttcagtga 396

Scoring table: OLIGO_NUC

Gapop 60.0 , Gapext 60.0

Searched: 441362 seqs, 153338381 residues

d size : 0

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : Issued_Patents_NA:*

1: /cgn2_6/ptodata/1/ina/5A_COMB.seq:*
2: /cgn2_6/ptodata/1/ina/5B_COMB.seq:*
3: /cgn2_6/ptodata/1/ina/6A_COMB.seq:*
4: /cgn2_6/ptodata/1/ina/6B_COMB.seq:*
5: /cgn2_6/ptodata/1/ina/PCRTUS_COMB.seq:*
6: /cgn2_6/ptodata/1/ina/Backfillseq1.seq:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	18	4.5	31	1	US-08-066-281-1
2	18	4.5	430	4	US-09-397-787-254
3	17	4.3	1118	4	US-09-452-239-37
4	17	4.3	1146	4	US-09-452-239-3
5	17	4.3	1381	4	US-08-858-207A-108
6	17	4.3	1509	1	US-08-115-052-1
7	17	4.3	2686	4	US-09-228-986-3
8	17	4.3	11613	4	US-09-453-702B-42
9	17	4.3	40328	3	US-08-742-185-102
10	16	4.0	36	4	US-08-910-722-7
11	16	4.0	42	4	US-08-910-722-5
12	16	4.0	57	1	US-08-474-177-17
13	16	4.0	57	1	US-08-487-033-17
14	16	4.0	57	1	US-08-480-810-17
15	16	4.0	57	2	US-08-508-735-17
16	16	4.0	57	2	US-08-848-251-17
17	16	4.0	57	2	US-08-848-047-17
18	16	4.0	57	3	US-09-120-130-17
19	16	4.0	57	3	US-09-115-252-17
20	16	4.0	57	3	US-08-986-515-17
21	16	4.0	57	3	US-09-120-128-17
22	16	4.0	57	4	US-09-120-129-17
23	16	4.0	57	4	US-09-201-139-17
24	16	4.0	57	4	US-09-120-131-17
25	16	4.0	384	4	US-09-134-001C-1752
26	16	4.0	471	1	US-08-474-177-1
27	16	4.0	471	1	US-08-487-033-1

C 28	16	4.0	471	1	US-08-480-810-1	Sequence 1, Appli
C 29	16	4.0	471	2	US-08-508-735-1	Sequence 1, Appli
C 30	16	4.0	471	2	US-08-848-251-1	Sequence 1, Appli
C 31	16	4.0	471	2	US-08-486-047-1	Sequence 1, Appli
C 32	16	4.0	471	3	US-09-120-130-1	Sequence 1, Appli
C 33	16	4.0	471	3	US-09-115-252-1	Sequence 1, Appli
C 34	16	4.0	471	3	US-08-986-515-1	Sequence 1, Appli
C 35	16	4.0	471	3	US-09-120-128-1	Sequence 1, Appli
C 36	16	4.0	471	4	US-09-120-129-1	Sequence 1, Appli
C 37	16	4.0	471	4	US-09-201-139-1	Sequence 1, Appli
C 38	16	4.0	471	4	US-09-120-131-1	Sequence 1, Appli
C 39	16	4.0	536	4	US-09-221-017B-1081	Sequence 1081, Ap
C 40	16	4.0	687	4	US-09-457-568-23	Sequence 23, Appl
C 41	16	4.0	687	4	US-09-457-646-23	Sequence 23, Appl
C 42	16	4.0	737	4	US-09-457-568-19	Sequence 19, Appl
C 43	16	4.0	737	4	US-09-457-646-19	Sequence 19, Appl
C 44	16	4.0	782	4	US-09-457-568-21	Sequence 21, Appl
C 45	16	4.0	782	4	US-09-457-646-21	Sequence 21, Appl

ALIGNMENTS

```
RESULT 1
US-08-066-281-1
; Sequence 1, Application US/08066281
; Patent No. 5376533
; GENERAL INFORMATION:
; APPLICANT: MacIaren, No. 53765331 K.
; APPLICANT: Song, Yao Hua
; TITLE OF INVENTION: Methods and Compositions for the Detection of
; TITLE OF INVENTION: Addison's Disease
; NUMBER OF SEQUENCES: 12
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: DAVID R. SALIWANCHIK
; STREET: 2421 N.W. 41st Street, Suite A-1
; CITY: Gainesville
; STATE: FL
; COUNTRY: USA
; ZIP: 32606
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentln Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/066,281
; FILING DATE: 19930524
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: SalIWanchik, David R.
; REGISTRATION NUMBER: 31,794
; REFERENCE/DOCKET NUMBER: UF/S&S-132
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 904-375-8100
; TELEFAX: 904-372-5800
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 31 bases
; TYPE: NUCLEIC ACID
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (synthetic)
; US-08-066-281-1

Query Match 4.5%; Score 18; DB 1; Length 31;
Best Local Similarity 100.0%; Pred. No. 10;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 288 TCCATGCTGCTCTCTGGGC 305
|||||
Db 8 TCCATGCTGCTCTCTGGGC 25
```

```
RESULT 2
US-09-397-787-254/C
: Sequence 254, Application US/09397787
: Patent No. 6468758
: GENERAL INFORMATION:
: APPLICANT: Benson, Darin R.
: APPLICANT: Lodes, Michael J.
: APPLICANT: Mitcham, Jennifer L.
: APPLICANT: King, Gordon E.
: TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR OVARIAN
: FILE REFERENCE: 210121.466C2
: CURRENT APPLICATION NUMBER: US/09/397,787
: CURRENT FILING DATE: 1999-09-16
: NUMBER OF SEQ ID NOS: 334
: SOFTWARE: FastSeq for Windows Version 3.0
: SEQ ID NO 254
: LENGTH: 430
: TYPE: DNA
: ORGANISM: Homo sapien
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (1)..(430)
: OTHER INFORMATION: n = A,T,C or G
US-09-397-787-254

Query Match          4.5%; Score 18; DB 4; Length 430;
Best Local Similarity 100.0%; Pred. No. 9.4;
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 191 GCGTTGCAAGATCTGCG 208
DB 273 GCGTTGCAAGATCTGCG 256

RESULT 3
US-09-452-239-37
: Sequence 37, Application US/09452239
: Patent No. 6465229
: GENERAL INFORMATION:
: APPLICANT: Rafalski, Antoni J.
: APPLICANT: Fader, Gary M.
: APPLICANT: Cahoon, Rebecca E.
: TITLE OF INVENTION: Plant Caffey1-CoA O-Methyltransferase
: FILE REFERENCE: B81284 US NA
: CURRENT APPLICATION NUMBER: US/09/452,239
: CURRENT FILING DATE: 1999-12-01
: EARLIER APPLICATION NUMBER: 60/110,594
: EARLIER FILING DATE: 1998-December-02
: NUMBER OF SEQ ID NOS: 50
: SOFTWARE: Microsoft Office 97
: SEQ ID NO 37
: LENGTH: 1118
: TYPE: DNA
: ORGANISM: Trifolium aestivum
US-09-452-239-37

Query Match          4.3%; Score 17; DB 4; Length 1118;
Best Local Similarity 100.0%; Pred. No. 28;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 273 GCGTTTTTTTGGCTC 289
DB 971 GCGTTTTTTTGGCTC 987

RESULT 4
US-09-452-239-3
: Sequence 3, Application US/09452239
: Patent No. 6465229
: GENERAL INFORMATION:
: APPLICANT: Rafalski, Antoni J.
```

```
: APPLICANT: Fader, Gary M.
: APPLICANT: Cahoon, Rebecca E.
: TITLE OF INVENTION: Plant Caffey1-CoA O-Methyltransferase
: FILE REFERENCE: B81284 US NA
: CURRENT APPLICATION NUMBER: US/09/452,239
: CURRENT FILING DATE: 1999-12-01
: EARLIER APPLICATION NUMBER: 60/110,594
: EARLIER FILING DATE: 1998-December-02
: NUMBER OF SEQ ID NOS: 50
: SOFTWARE: Microsoft Office 97
: SEQ ID NO 3
: LENGTH: 1146
: TYPE: DNA
: ORGANISM: Zea mays
US-09-452-239-3

Query Match          4.3%; Score 17; DB 4; Length 1146;
Best Local Similarity 100.0%; Pred. No. 28;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 273 GCGTTTTTTTGGCTC 289
DB 976 GCGTTTTTTTGGCTC 992

RESULT 5
US-08-858-207A-108/C
: Sequence 108, Application US/08858207A
: Patent No. 6348328
: GENERAL INFORMATION:
: APPLICANT: Black, Michael
: APPLICANT: Hodgson, John
: APPLICANT: Knowles, David
: APPLICANT: Nicholas, Richard
: APPLICANT: Stodola, Robert
: TITLE OF INVENTION: NO. 6348328e1 Compounds
: NUMBER OF SEQUENCES: 552
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: SmithKline Beecham Corporation
: STREET: 709 Swedeland Road
: CITY: King of Prussia
: STATE: PA
: COUNTRY: USA
: ZIP: 19406-0939
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Diskette
: COMPUTER: IBM Compatible
: OPERATING SYSTEM: DOS
: SOFTWARE: FastSeq for Windows Version 2.0
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/858,207A
: FILING DATE: 09-MAY-1997
: CLASSIFICATION: 435
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: 60/017670
: FILING DATE: 14-MAY-1996
: ATTORNEY/AGENT INFORMATION:
: NAME: Gimml, Edward R
: REGISTRATION NUMBER: 38,891
: REFERENCE/DOCKET NUMBER: P50475
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: 610-270-4478
: TELEFAX: 610-270-5090
: TELEX:
: INFORMATION FOR SEQ ID NO: 108:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 1381 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
US-08-858-207A-108

Query Match          4.3%; Score 17; DB 4; Length 1381;
```

Best Local Similarity 100.0%; Pred. No. 28;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 323 CATTTCTCTCATCTC 339
|||||
Db 295 CATTTCTCTCATCTC 279

RESULT 6

US-08-115-052-1
; Sequence 1, Application US/08115052
; Patent No. 5705400
; GENERAL INFORMATION:
; APPLICANT: Furmaniak-Wehr, Jadwiga Maria
; TITLE OF INVENTION: Assay for Adrenal Autoantigen
; NUMBER OF SEQUENCES: 2
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Banner, Birch, McKie & Beckett
; STREET: 1001 G Street N.W.
; CITY: Washington
; STATE: D.C.
; COUNTRY: USA
; ZIP: 20001
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/115.052
; FILING DATE: 02-SEP-1993
; CLASSIFICATION: 424
; PRIOR APPLICATION NUMBER: 07/937,409
; APPLICATION NUMBER: 31-AUG-1992
; FILING DATE: 31-AUG-1992
; ATTORNEY/AGENT INFORMATION:
; NAME: Dale H. Hoscheit
; REGISTRATION NUMBER: 19,090
; REFERENCE/DOCKET NUMBER: 01950.44179
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-508-9100
; TELEFAX: 202-5089299
; TELEX: 197430 BBMB UT
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1509 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: CDNA
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; DEVELOPMENTAL STAGE: foetus
; TISSUE TYPE: adrenal gland
; FEATURE:
; NAME/KEY: sig_peptide
; LOCATION: 13..54
; NAME/KEY: mat_peptide
; LOCATION: 55..1494
; OTHER INFORMATION: /product="steroid 21-hydroxylase"
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 13..1494
; NAME/KEY: misc_feature
; LOCATION: (435-436)
; OTHER INFORMATION: /standard_name="PvuII cleavage
; OTHER INFORMATION: site"
; FEATURE:
; NAME/KEY: misc_feature

LOCATION: (732-733)
; OTHER INFORMATION: /standard_name="PvuII cleavage
; OTHER INFORMATION: site"
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (852-853)
; OTHER INFORMATION: /standard_name="PnaCI cleavage
; OTHER INFORMATION: site"
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1148-1149)
; OTHER INFORMATION: /standard_name="SauI cleavage
; OTHER INFORMATION: site"
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1354-1355)
; OTHER INFORMATION: /standard_name="StuI cleavage
; OTHER INFORMATION: site"
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (53-54)
; OTHER INFORMATION: /standard_name="NarI cleavage
; OTHER INFORMATION: site"

US-08-115-052-1

Query Match 4.3%; Score 17; DB 1; Length 1509;
Best Local Similarity 100.0%; Pred. No. 28;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 289 CCATGCTGCTCCTGGC 305
|||||
Db 11 CCATGCTGCTCCTGGC 27

RESULT 7

US-09-228-986-3
; Sequence 3, Application US/09228986
; Patent No. 6359198
; GENERAL INFORMATION:
; APPLICANT: Strabala, Timothy
; APPLICANT: Nieuwenhuizen, Niels
; TITLE OF INVENTION: Compositions Isolated from Plant Cells
; FILE REFERENCE: 11000/1020
; CURRENT APPLICATION NUMBER: US/09/228,986
; CURRENT FILING DATE: 1999-01-12
; NUMBER OF SEQ. ID NOS: 130
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 3
; LENGTH: 2686
; TYPE: DNA
; ORGANISM: Pinus radiata
; US-09-228-986-3

Query Match 4.3%; Score 17; DB 4; Length 2686;
Best Local Similarity 100.0%; Pred. No. 27;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 318 TACCCATTTTCTCA 334
|||||
Db 272 TACCCATTTTCTCA 288

RESULT 8

US-09-453-702B-42/c
; Sequence 42, Application US/09453702B
; Patent No. 6365723
; GENERAL INFORMATION:
; APPLICANT: Blattner, Frederick R.
; Blattner, Valerie
; Perna, Nicole T.
; Plunkett, Guy
; Welch, Rod

TITLE OF INVENTION: No. 6365723el Sequences of E. coli 0157
NUMBER OF SEQUENCES: 265
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Quarles & Brady
STREET: 1 South Plinckney Street
CITY: Madison
STATE: WI
COUNTRY: US
ZIP: 53701-2113
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette, 3.50 inch, 1.44mb storage
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Word Perfect 8.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/453,702B
FILING DATE: 03-Dec-1999
CLASSIFICATION: <unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 60/110,955
FILING DATE: 04-DEC-1998
ATTORNEY/AGENT INFORMATION:
NAME: Seay, Nicholas J.
REGISTRATION NUMBER: 27386
REFERENCE/DOCKET NUMBER: 960296.95017
TELECOMMUNICATION INFORMATION:
TELEPHONE: (608) 251-5000
TELEFAX: (608) 251-9166
INFORMATION FOR SEQ ID NO: 42:
SEQUENCE CHARACTERISTICS:
LENGTH: 11613
TYPE: nucleic acid
STRANDEDNESS: double
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
SEQUENCE DESCRIPTION: SEQ ID NO: 42:
US-09-453-702B-42

Query Match 4.3%; Score 17; DB 4; Length 11613;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 219 GGTTCCTGATGGGCT 235
|||||
DB 2106 GGTTCCTGATGGGCT 2090

RESULT 9
Sequence 102, Application US/08742185
Patent No. 6020476
GENERAL INFORMATION:
APPLICANT: Page, David C.
APPLICANT: Reljo, Renee
APPLICANT: Saxena, Richa
APPLICANT: Hawkins, Trevor
APPLICANT: Reene, Mary Pat
TITLE OF INVENTION: DAZ: A GENE FAMILY ASSOCIATED WITH AZOOSPERMIA
NUMBER OF SEQUENCES: 102
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Hamilton, Brook, Smith & Reynolds, P.C.
STREET: Two Militia Drive
CITY: Lexington
STATE: Massachusetts
COUNTRY: US
ZIP: 02173
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/742,185

FILING DATE: 30-OCT-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/690,734
FILING DATE: 31-JUL-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/310,429
FILING DATE: 22-SEP-1994
ATTORNEY/AGENT INFORMATION:
NAME: Granahan, Patricia
REGISTRATION NUMBER: 32,227
REFERENCE/DOCKET NUMBER: WH194-07A2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (617) 861-6240
TELEFAX: (617) 861-9540
INFORMATION FOR SEQ ID NO: 102:
SEQUENCE CHARACTERISTICS:
LENGTH: 40328 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
US-08-742-185-102

Query Match 4.3%; Score 17; DB 3; Length 40328;
Best Local Similarity 100.0%; Pred. No. 26;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 268 GGTGGGCTTTT TTTT 284
|||||
DB 882 GGTGGGCTTTT TTTT 898

RESULT 10
US-08-910-722-7/c
Sequence 77, Application US/08910722
Patent No. 6251871
GENERAL INFORMATION:
APPLICANT: Jin, Xiaomei
APPLICANT: Roth, Jack A.
TITLE OF INVENTION: p16 EXPRESSION CONSTRUCTS AND THEIR
TITLE OF INVENTION: APPLICATION IN CANCER THERAPY
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Arnold, White & Durkee
STREET: P.O. Box 4433
CITY: Houston
STATE: Texas
COUNTRY: United States of America
ZIP: 77210
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
OPERATING SYSTEM: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/910,722
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/502,881
FILING DATE:
ATTORNEY/AGENT INFORMATION:
NAME: Highlander, Steven L.
REGISTRATION NUMBER: 37,642
REFERENCE/DOCKET NUMBER: INGN.016/HYL
TELECOMMUNICATION INFORMATION:
TELEPHONE: (512) 418-3000
TELEFAX: (512) 474-7577
TELE: 79-0924
INFORMATION FOR SEQ ID NO: 7:
SEQUENCE CHARACTERISTICS:
LENGTH: 36 base pairs

TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "DNA"
US-08-910-722-7

Query Match 4.0%; Score 16; DB 4; Length 36;
Best Local Similarity 100.0%; Pred. No. 93;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 285 GGCTCCATGCTGCTCC 300
|||||
DB 23 GGCTCCATGCTGCTCC 8

RESULT 11
US-08-910-722-5/C
Sequence 5, Application US/08910722
Patent No. 6251871

GENERAL INFORMATION:

APPLICANT: Jin, Xiaomei
APPLICANT: Roth, Jack A.
TITLE OF INVENTION: p16 EXPRESSION CONSTRUCTS AND THEIR
TITLE OF INVENTION: APPLICATION IN CANCER THERAPY
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESS:
ADDRESSEE: Arnold, White & Durkee
STREET: P.O. Box 4433
CITY: Houston
STATE: Texas
COUNTRY: United States of America
ZIP: 77210
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/910,722
FILING DATE:

CLASSIFICATION:

PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/502,881
FILING DATE:

ATTORNEY/AGENT INFORMATION:

NAME: Highlander, Steven L.
REGISTRATION NUMBER: 37,642
REFERENCE/DOCKET NUMBER: INGN:016/HYL
TELECOMMUNICATION INFORMATION:
TELEPHONE: (512) 418-3000
TELEFAX: (512) 474-7577
TELEX: 79-0924

INFORMATION FOR SEQ ID NO: 5:

SEQUENCE CHARACTERISTICS:
LENGTH: 42 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: other nucleic acid
DESCRIPTION: /desc = "DNA"
US-08-910-722-5

Query Match 4.0%; Score 16; DB 4; Length 42;
Best Local Similarity 100.0%; Pred. No. 93;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 285 GGCTCCATGCTGCTCC 300
|||||
DB 29 GGCTCCATGCTGCTCC 14

RESULT 12

US-08-474-177-17/C
Sequence 17, Application US/08474177
Patent No. 5624819

GENERAL INFORMATION:

APPLICANT: Skolnick, Mark H.
APPLICANT: Cannon-Albright, Lisa A.
TITLE OF INVENTION: GERMLINE MUTATIONS IN THE MTS GENE
NUMBER OF SEQUENCES: 36
CORRESPONDENCE ADDRESS:
ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
STREET: 1201 New York Avenue, Suite 1000
CITY: Washington
STATE: DC
COUNTRY: USA
ZIP: 20005

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/474,177
FILING DATE: 07-JUN-1995
CLASSIFICATION: 435

PRIOR APPLICATION DATA:

APPLICATION NUMBER: PCT/US95/03537
FILING DATE: 17-MAR-1995

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/251,938
FILING DATE: 01-JUN-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/215,087
FILING DATE: 18-MAR-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/215,086
FILING DATE: 18-MAR-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/227,369
FILING DATE: 14-APR-1994

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/214,582
FILING DATE: 18-MAR-1994

ATTORNEY/AGENT INFORMATION:

NAME: Ihnen, Jeffrey L.
REGISTRATION NUMBER: 28,957
REFERENCE/DOCKET NUMBER: 24884-109348-E
TELECOMMUNICATION INFORMATION:
TELEPHONE: 202-962-4810

INFORMATION FOR SEQ ID NO: 17:

SEQUENCE CHARACTERISTICS:
LENGTH: 57 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHEICAL: NO
ANTI-SENSE: NO
ORIGINAL SOURCE:
ORGANISM: Homo sapiens
US-08-474-177-17

Query Match 4.0%; Score 16; DB 1; Length 57;
Best Local Similarity 100.0%; Pred. No. 92;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 285 GGCTCCATGCTGCTCC 300
|||||
DB 50 GGCTCCATGCTGCTCC 35

RESULT 13

US-08-487-033-17/c
; Sequence 17, Application US/08487033
; Patent No. 5739027
; GENERAL INFORMATION:
; APPLICANT: Kamb, Alexander
; TITLE OF INVENTION: MTS1-Beta GENE
; NUMBER OF SEQUENCES: 36
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
; STREET: 1201 New York Avenue, Suite 1000
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/487,033
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/03316
; FILING DATE: 17-MAR-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/251,938
; FILING DATE: 01-JUN-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/215,087
; FILING DATE: 18-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/215,086
; FILING DATE: 18-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/227,369
; FILING DATE: 14-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/214,582
; FILING DATE: 18-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ihnen, Jeffrey L.
; REGISTRATION NUMBER: 28,957
; REFERENCE/DOCKET NUMBER: 24884-109348-C
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-962-8300
; TELEFAX: 202-962-8300
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 57 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; US-08-487-033-17

Query Match 4.0%; Score 16; DB 1; Length 57;
Best Local Similarity 100.0%; Pred. No. 92;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 285 GGCTCATGCTGCTCC 300
|||||
Db 50 GGCTCATGCTGCTCC 35

RESULT 14
US-08-480-810-17/c
; Sequence 17, Application US/08480810

; Patent No. 5801236
; GENERAL INFORMATION:
; APPLICANT: Kamb, Alexander
; TITLE OF INVENTION: MTS1 GENE
; NUMBER OF SEQUENCES: 36
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
; STREET: 1201 New York Avenue, Suite 1000
; CITY: Washington
; STATE: DC
; COUNTRY: USA
; ZIP: 20005
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/480,810
; FILING DATE: 07-JUN-1995
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/03316
; FILING DATE: 17-MAR-1995
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/251,938
; FILING DATE: 01-JUN-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/215,087
; FILING DATE: 18-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/215,086
; FILING DATE: 18-MAR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/227,369
; FILING DATE: 14-APR-1994
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/214,582
; FILING DATE: 18-MAR-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Ihnen, Jeffrey L.
; REGISTRATION NUMBER: 28,957
; REFERENCE/DOCKET NUMBER: 24884-109348
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 202-962-8300
; TELEFAX: 202-962-8300
; INFORMATION FOR SEQ ID NO: 17:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 57 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; ORIGINAL SOURCE:
; ORGANISM: Homo sapiens
; US-08-480-810-17

Query Match 4.0%; Score 16; DB 1; Length 57;
Best Local Similarity 100.0%; Pred. No. 92;
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 285 GGCTCATGCTGCTCC 300
|||||
Db 50 GGCTCATGCTGCTCC 35

RESULT 15
US-08-508-735-17/c
; Sequence 17, Application US/08508735
; Patent No. 5843756
; GENERAL INFORMATION:

APPLICANT: Stone, Steven
 APPLICANT: Jang, Ping
 APPLICANT: Kamb, Alexander
 TITLE OF INVENTION: MTS GENE AND THERAPEUTIC USE THEREOF
 NUMBER OF SEQUENCES: 47
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Venable, Baetjer, Howard & Civiletti, LLP
 STREET: 1201 New York Avenue, Suite 1000
 CITY: Washington
 STATE: DC
 COUNTRY: USA
 ZIP: 20005
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: Patent Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/508,735
 FILING DATE:
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US to be assigned
 FILING DATE: 07-JUN-1995
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: PCT/US95/03316
 FILING DATE: 17-MAR-1995
 ATTORNEY/AGENT INFORMATION:
 NAME: Ihnen, Jeffrey L.
 REGISTRATION NUMBER: 28,957
 REFERENCE/DOCKET NUMBER: 24884-109348
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 202-962-8300
 TELEFAX: 202-962-4848
 INFORMATION FOR SEQ ID NO: 17:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 57 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 HYPOTHETICAL: NO
 ANTI-SENSE: NO
 ORIGINAL SOURCE:
 ORGANISM: Homo sapiens
 US-08-508-735-17

Query Match 4.0%; Score 16; DB 2; Length 57;
 Best Local Similarity 100.0%; Pred. NO. 92;
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 285 GGCTCCATGCTGCTCC 300
 ||||||||||||
 Db 50 GGCTCCATGCTGCTCC 35

Search completed: April 25, 2003, 00:54:16
 Job time : 56.8639 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 00:41:49 ; Search time 81.2308 Seconds
(without alignments)
5304.628 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_446

Perfect score: 396
Sequence: 1 atgggtgacatttccttgcctt.....gamctgatacttcagtga 396

Scoring table:
OLIGO_NUC
Gapop 60.0 , Gapext 60.0

Searched: 709820 seqs, 544064369 residues

d size : 0

Total number of hits satisfying chosen parameters: 1419640

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Listing first 45 summaries

Database : Published_Applications_NA:*

1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq:*
2: /cgn2_6/ptodata/1/pubpna/PC1_NEW_PUB.seq:*
3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq:*
4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq:*
5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq:*
6: /cgn2_6/ptodata/1/pubpna/PC105_PUBCOMB.seq:*
7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq:*
8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq:*
9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq:*
11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*
12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq:*
13: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
14: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	33	8.3	684973	10 US-09-263-959-1	Sequence 1, Appli
C 2	22	5.6	361	9 US-09-918-995-8549	Sequence 8549, Ap
C 3	19	4.8	802	9 US-09-974-879-27	Sequence 27, Appl
C 4	18	4.5	430	10 US-09-876-889-254	Sequence 254, Appl
C 5	18	4.5	431	10 US-09-866-562-80	Sequence 80, Appl
C 6	18	4.5	431	10 US-09-866-562-87	Sequence 87, Appl
C 7	18	4.5	570	10 US-09-864-761-9118	Sequence 9118, Ap
C 8	17	4.3	489	9 US-09-918-995-28049	Sequence 28049, A
C 9	17	4.3	491	9 US-09-918-995-34650	Sequence 34650, A
C 10	17	4.3	492	9 US-09-918-995-19262	Sequence 19262, A
C 11	17	4.3	531	9 US-10-092-154-1899	Sequence 1899, Ap
C 12	17	4.3	553	10 US-09-764-847-1899	Sequence 1899, Ap
C 13	17	4.3	553	10 US-09-864-761-13668	Sequence 13668, A
C 14	17	4.3	609	10 US-09-974-300-8084	Sequence 8084, Ap
C 15	17	4.3	944	9 US-09-774-639-99	Sequence 99, Appl
C 16	17	4.3	944	9 US-09-969-730-16	Sequence 16, Appl
C 17	17	4.3	1047	10 US-09-823-830A-468	Sequence 468, App
C 18	17	4.3	1089	10 US-09-962-740-3	Sequence 3, Appli
C 19	17	4.3	1118	10 US-09-452-239-37	Sequence 37, Appli

20	17	4.3	1146	10 US-09-452-239-3	Sequence 3, Appli
C 21	17	4.3	1215	10 US-09-962-740-1	Sequence 1, Appli
C 22	17	4.3	1215	10 US-09-962-740-6	Sequence 6, Appli
C 23	17	4.3	1396	10 US-09-962-740-8	Sequence 8, Appli
C 24	17	4.3	1631	10 US-09-962-740-5	Sequence 5, Appli
C 25	17	4.3	2686	9 US-10-101-464A-3	Sequence 3, Appli
C 26	17	4.3	2880	9 US-09-951-502A-1	Sequence 1, Appli
C 27	17	4.3	11613	9 US-10-114-170-42	Sequence 42, Appli
C 28	17	4.3	38374	10 US-09-880-107-3463	Sequence 3463, Ap
C 29	17	4.3	172637	10 US-09-805-458A-3	Sequence 3, Appli
C 30	17	4.3	326014	10 US-09-731-231A-3	Sequence 3, Appli
C 31	17	4.3	1503841	9 US-09-946-807-1	Sequence 1, Appli
C 32	17	4.3	1503841	10 US-09-795-668-1	Sequence 1, Appli
C 33	17	4.0	121	9 US-09-818-875-1331	Sequence 1321, Ap
C 34	16	4.0	121	9 US-09-818-875-1332	Sequence 1325, Ap
C 35	16	4.0	121	9 US-09-818-875-1335	Sequence 1329, Ap
C 36	16	4.0	121	9 US-09-818-875-1336	Sequence 1330, Ap
C 37	16	4.0	121	9 US-09-818-875-1337	Sequence 1333, Ap
C 38	16	4.0	121	9 US-09-818-875-1339	Sequence 1334, Ap
C 39	16	4.0	121	9 US-09-818-875-1340	Sequence 1334, Ap
C 40	16	4.0	121	9 US-09-818-875-1341	Sequence 1334, Ap
C 41	16	4.0	121	9 US-09-818-875-1342	Sequence 1334, Ap
C 42	16	4.0	235	10 US-09-878-574-13480	Sequence 5259, Ap
C 43	16	4.0	281	10 US-09-878-574-13480	Sequence 13480, A
C 44	16	4.0	281	10 US-09-964-824A-385	Sequence 385, App
C 45	16	4.0	321	10 US-09-864-761-25530	Sequence 25530, A

ALIGNMENTS

RESULT 1
US-09-263-959-1/c
Sequence 1, Application US/09263959
Patent No. US20020150891A1
GENERAL INFORMATION:
APPLICANT: Hood, Leroy E.
APPLICANT: Kowen, Ben F.
TITLE OF INVENTION: DIAGNOSTIC AND THERAPEUTIC COMPOSITIONS AND METHODS WHICH U
NUMBER OF SEQUENCES: 1279
CORRESPONDENCE ADDRESS:
ADDRESSEE: Seed and Berry LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington
COUNTRY: US
ZIP: 98104-7092
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patentin Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/263,959
FILING DATE: 05-MAR-1999
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION:
NAME: Mcmasters, David D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 920010.426C2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (206) 622-4900
TELEFAX: (206) 682-6031
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 684973 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
US-09-263-959-1
Query Match 8.3%, Score 33, DB 10, Length 684973;

Best Local Similarity 100.0%; Pred. No. 6,4e-08;
Matches 33: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 66 TGTAAACTCCGGCTCTGTGTGCTGAG 98
|||||
Db 404690 TGTAAACTCCTGGCTCTGTGTGCTGAG 404658

RESULT 2

US-09-918-995-8549/c
; Sequence 8549, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; PRIOR FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: PastSeq for Windows Version 3.0
; SEQ ID NO 8549
; LENGTH: 361
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-918-995-8549

Query Match
Best Local Similarity 100.0%; Pred. No. 0.063;
Matches 22: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 149 GGCCTGTGTCATGGCTCAG 170
|||||
Db 143 GGCCTGTGTCATGGCTCAG 122

RESULT 3

US-09-974-879-27/c
; Sequence 27, Application US/09974879
; Publication No. US20030028003A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: 125 Human Secreted Proteins
; FILE REFERENCE: P2020P2
; CURRENT APPLICATION NUMBER: US/09/974,879
; PRIOR FILING DATE: 2001-10-12
; PRIOR APPLICATION NUMBER: US 60/239,893
; PRIOR FILING DATE: 2000-10-13
; PRIOR APPLICATION NUMBER: US 09/818,683
; PRIOR FILING DATE: 2001-03-28
; PRIOR APPLICATION NUMBER: US 09/305,736
; PRIOR FILING DATE: 1999-05-05
; PRIOR APPLICATION NUMBER: PCT/US98/23435
; PRIOR FILING DATE: 1998-11-04
; PRIOR APPLICATION NUMBER: US 60/064,911
; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/064,912
; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/064,983
; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/064,900
; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/064,988
; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/064,987
; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/064,908
; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/064,984
; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/064,985

; PRIOR FILING DATE: 1997-11-07
; PRIOR APPLICATION NUMBER: US 60/066,094

; PRIOR FILING DATE: 1997-11-17
; PRIOR APPLICATION NUMBER: US 60/066,100
; PRIOR FILING DATE: 1997-11-17
; PRIOR APPLICATION NUMBER: US 60/066,089
; PRIOR FILING DATE: 1997-11-17
; PRIOR APPLICATION NUMBER: US 60/066,095
; PRIOR FILING DATE: 1997-11-17
; PRIOR APPLICATION NUMBER: US 60/066,090
; PRIOR FILING DATE: 1997-11-17
; NUMBER OF SEQ ID NOS: 611
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 27

; LENGTH: 802
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: SITE
; LOCATION: (337)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: SITE
; LOCATION: (359)
; OTHER INFORMATION: n equals a,t,g, or c
US-09-974-879-27

Query Match
Best Local Similarity 100.0%; Pred. No. 2.5;
Matches 19: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 217 GGTGTTCTGATGGGT 235
|||||
Db 304 GTGTTCTGATGGGT 286

RESULT 4

US-09-876-889-254/c
; Sequence 254, Application US/09876889
; Patent No. US20020076715A1
; GENERAL INFORMATION:
; APPLICANT: Benson, Darin R.
; APPLICANT: Lodes, Michael J.
; APPLICANT: Mitcham, Jennifer L.
; APPLICANT: King, Gordon E.
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR OVARIAN
; FILE REFERENCE: 210121.466C3
; CURRENT APPLICATION NUMBER: US/09/876,889
; CURRENT FILING DATE: 2001-06-06
; NUMBER OF SEQ ID NOS: 353
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 254
; LENGTH: 430
; TYPE: DNA
; ORGANISM: Homo sapien
; FEATURE:
; NAME/KEY: misc.feature
; LOCATION: (1)-(430)
; OTHER INFORMATION: n = A,T,C or G
US-09-876-889-254

Query Match
Best Local Similarity 100.0%; Pred. No. 8.7;
Matches 18: Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 191 GGGTGCAGATCTGTG 208
|||||
Db 273 GGGTGCAGATCTGTG 256

RESULT 5

US-09-866-562-80
; Sequence 80, Application US/09866562

Patent No. US20020009758A1
 GENERAL INFORMATION:
 APPLICANT: Harlocker, Susan L.
 APPLICANT: Wang, Tongtong
 APPLICANT: Bangur, Chaitanya S.
 APPLICANT: Klee, Jennifer
 APPLICANT: Switzer, Anne
 TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
 TITLE OF INVENTION: AND DIAGNOSIS OF LUNG CANCER.
 FILE REFERENCE: 210121.502
 CURRENT APPLICATION NUMBER: US/09/866,562
 CURRENT FILING DATE: 2001-05-25
 NUMBER OF SEQ ID NOS: 96
 SEQ ID NO 80
 LENGTH: 431
 TYPE: DNA
 ORGANISM: Homo sapiens
 FEATURE:
 NAME/KEY: misc-feature
 LOCATION: 361..431
 OTHER INFORMATION: n = A,T,C or G
 09-866-562-80

Query Match 4.5%; Score 18; DB 10; Length 431;
 Best Local Similarity 100.0%; Pred. No. 8.7;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 91 TGCTGAGTGGCTGCTCT 108
 ||||||||||||||||
 Db 18 TGCTGAGTGGCTGCTCT 35

RESULT 6
 US-09-866-562-87
 Sequence 87, Application US/09866562
 Patent No. US20020009758A1
 GENERAL INFORMATION:
 APPLICANT: Harlocker, Susan L.
 APPLICANT: Wang, Tongtong
 APPLICANT: Bangur, Chaitanya S.
 APPLICANT: Klee, Jennifer
 APPLICANT: Switzer, Anne
 TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY
 TITLE OF INVENTION: AND DIAGNOSIS OF LUNG CANCER.
 FILE REFERENCE: 210121.502
 CURRENT APPLICATION NUMBER: US/09/866,562
 CURRENT FILING DATE: 2001-05-25
 NUMBER OF SEQ ID NOS: 96
 SEQ ID NO 87
 LENGTH: 431
 TYPE: DNA
 ORGANISM: Homo sapiens
 FEATURE:
 NAME/KEY: misc-feature
 LOCATION: 361..431
 OTHER INFORMATION: n = A,T,C or G
 US-09-866-562-87

Query Match 4.5%; Score 18; DB 10; Length 431;
 Best Local Similarity 100.0%; Pred. No. 8.7;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 91 TGCTGAGTGGCTGCTCT 108
 ||||||||||||||||
 Db 18 TGCTGAGTGGCTGCTCT 35
 RESULT 7
 US-09-864-761-9118/c
 Sequence 9118, Application US/09864761
 Patent No. US20020048763A1
 GENERAL INFORMATION:
 APPLICANT: Penn, Sharon G.

APPLICANT: Rank, David R.
 APPLICANT: Hanzel, David K.
 APPLICANT: Chen, Wensheng
 TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FO
 FILE REFERENCE: Aeonica-X-1
 CURRENT APPLICATION NUMBER: US/09/864,761
 CURRENT FILING DATE: 2001-05-23
 PRIOR APPLICATION NUMBER: US 60/180,312
 PRIOR FILING DATE: 2000-02-04
 PRIOR APPLICATION NUMBER: US 60/207,456
 PRIOR FILING DATE: 2000-05-26
 PRIOR APPLICATION NUMBER: US 09/632,366
 PRIOR FILING DATE: 2000-08-03
 PRIOR APPLICATION NUMBER: GB 24263.6
 PRIOR FILING DATE: 2000-10-04
 PRIOR APPLICATION NUMBER: US 60/236,359
 PRIOR FILING DATE: 2000-09-27
 PRIOR APPLICATION NUMBER: PCT/US01/00666
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00669
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00665
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00668
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00663
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00662
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00661
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: PCT/US01/00670
 PRIOR FILING DATE: 2001-01-30
 PRIOR APPLICATION NUMBER: US 60/234,687
 PRIOR FILING DATE: 2000-09-21
 PRIOR APPLICATION NUMBER: US 09/608,408
 PRIOR FILING DATE: 2000-06-30
 PRIOR APPLICATION NUMBER: US 09/774,203
 PRIOR FILING DATE: 2001-01-29
 NUMBER OF SEQ ID NOS: 49117
 SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
 SEQ ID NO 918
 LENGTH: 570
 TYPE: DNA
 ORGANISM: Homo sapiens
 FEATURE:
 OTHER INFORMATION: MAP TO AP000053.1
 OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.8
 OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 2.4
 OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 4.8
 OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 4
 OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 2.3
 OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 3.6
 OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 4.4
 US-09-864-761-9118

Query Match 4.5%; Score 18; DB 10; Length 570;
 Best Local Similarity 100.0%; Pred. No. 8.6;
 Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 85 TGTGTGCTGAGTGGC 102
 ||||||||||||||||
 Db 505 TGTGTGCTGAGTGGC 488

RESULT 8
 US-09-918-995-28049/c

```
; Sequence 28049, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 28049
; LENGTH: 489
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(489)
; OTHER INFORMATION: n = A,T,C or G
; US-918-995-28049
```

```
Query Match 4.3%; Score 17; DB 9; Length 489;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 325 TTTTCTTCATCTCTG 341
|||||
DB 416 TTTTCTTCATCTCTG 400
```

```
RESULT 9
US-09-918-995-34650/C
; Sequence 34650, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 34650
; LENGTH: 491
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(491)
; OTHER INFORMATION: n = A,T,C or G
; US-09-918-995-34650
```

```
Query Match 4.3%; Score 17; DB 9; Length 491;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 328 TTTCTCATCTCTGTGG 344
|||||
DB 138 TTTCTCATCTCTGTGG 122
```

```
RESULT 10
US-09-918-995-19262/C
; Sequence 19262, Application US/09918995
; Publication No. US20030073623A1
; GENERAL INFORMATION:
; APPLICANT: Hyseq, Inc.
; TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
```

```
; TITLE OF INVENTION: FROM VARIOUS CDNA LIBRARIES
; FILE REFERENCE: 20411-756
; CURRENT APPLICATION NUMBER: US/09/918,995
; CURRENT FILING DATE: 2001-07-30
; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 19262
; LENGTH: 492
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (1)..(492)
; OTHER INFORMATION: n = A,T,C or G
; US-09-918-995-19262
```

```
Query Match 4.3%; Score 17; DB 9; Length 492;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 246 CTCACGCTTCCTCTGG 262
|||||
DB 85 CTCACGCTTCCTCTGG 69
```

```
RESULT 11
US-10-092-154-1899/C
; Sequence 1899, Application US/10092154
; Publication No. US20030054575A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC009C1
; CURRENT APPLICATION NUMBER: US/10/092,154
; CURRENT FILING DATE: 2002-03-07
; NUMBER OF SEQ ID NOS: 2003
; PRIOR Application removed - See File Wrapper or Palm
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1899
; LENGTH: 531
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-10-092-154-1899
```

```
Query Match 4.3%; Score 17; DB 9; Length 531;
Best Local Similarity 100.0%; Pred. No. 30;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
```

```
OY 28 TCTTTTCATCTTGCA 44
|||||
DB 141 TCTTTTCATCTTGCA 125
```

```
RESULT 12
US-09-764-847-1899/C
; Sequence 1899, Application US/09764847
; Patent No. US20020132767A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PC009
; CURRENT APPLICATION NUMBER: US/09/764,847
; CURRENT FILING DATE: 2001-01-17
; PRIOR application data removed - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 2003
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1899
; LENGTH: 531
; TYPE: DNA
; ORGANISM: Homo sapiens
; US-09-764-847-1899
```


Query Match 4.3%; Score 17; DB 10; Length 531;
 Best Local Similarity 100.0%; Pred. No. 30;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 28 TCCTTTTCATCTTTGCA 44
 Db 141 TCCTTTTCATCTTTGCA 125

RESULT 13
 US-09-864-761-13668/c
 ; Sequence 13668, Application US/09864761
 ; Patent No. US2002048763A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Penn, Sharon G.
 ; APPLICANT: Rank, David R.
 ; APPLICANT: Hanzel, David K.
 ; APPLICANT: Chen, Wensheng
 ; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
 ; FILE REFERENCE: Aecomica-X-1
 ; CURRENT APPLICATION NUMBER: US/09/864,761
 ; PRIOR FILING DATE: 2001-05-23
 ; PRIOR APPLICATION NUMBER: US 60/180,312
 ; PRIOR FILING DATE: 2000-02-04
 ; PRIOR APPLICATION NUMBER: US 60/207,456
 ; PRIOR FILING DATE: 2000-05-26
 ; PRIOR APPLICATION NUMBER: US 09/632,366
 ; PRIOR FILING DATE: 2000-08-03
 ; PRIOR APPLICATION NUMBER: GB 24263.6
 ; PRIOR FILING DATE: 2000-10-04
 ; PRIOR APPLICATION NUMBER: US 60/236,359
 ; PRIOR FILING DATE: 2000-09-27
 ; PRIOR APPLICATION NUMBER: PCT/US01/00666
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00667
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00664
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00669
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00665
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00668
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00663
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00662
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00661
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: PCT/US01/00670
 ; PRIOR FILING DATE: 2001-01-30
 ; PRIOR APPLICATION NUMBER: US 60/234,687
 ; PRIOR FILING DATE: 2000-09-21
 ; PRIOR APPLICATION NUMBER: US 09/608,408
 ; PRIOR FILING DATE: 2000-06-30
 ; PRIOR APPLICATION NUMBER: US 09/774,203
 ; PRIOR FILING DATE: 2001-01-29
 ; SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
 ; SEQ ID NO 13668
 ; LENGTH: 553
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 ; FEATURE:
 ; OTHER INFORMATION: MAP TO 282201.1
 ; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.4
 US-09-864-761-13668

Query Match 4.3%; Score 17; DB 10; Length 553;
 Best Local Similarity 100.0%; Pred. No. 30; Mismatches 0; Indels 0; Gaps 0;

Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 327 TTCTTCATCTCTGCG 343
 Db 378 TTCTTCATCTCTGCG 362

RESULT 14
 US-09-974-300-8084/c
 ; Sequence 8084, Application US/09974300
 ; Patent No. US20020146721A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Berka, Randy M.
 ; APPLICANT: Clausen, Ib Groth
 ; TITLE OF INVENTION: Methods For Monitoring Multiple Gene
 ; FILE REFERENCE: 10085.500-US
 ; CURRENT APPLICATION NUMBER: US/09/974,300
 ; CURRENT FILING DATE: 2001-10-05
 ; PRIOR APPLICATION NUMBER: 09/680,598
 ; PRIOR FILING DATE: 2000-10-06
 ; PRIOR APPLICATION NUMBER: 60/279,526
 ; PRIOR FILING DATE: 2001-03-27
 ; NUMBER OF SEQ ID NOS: 8481
 ; SOFTWARE: FastSeq for Windows Version 4.0
 ; SEQ ID NO 8084
 ; LENGTH: 609
 ; TYPE: DNA
 ; ORGANISM: Bacillus clausii
 US-09-974-300-8084

Query Match 4.3%; Score 17; DB 10; Length 609;
 Best Local Similarity 100.0%; Pred. No. 30;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 28 TCCTTTTCATCTTTGCA 44
 Db 536 TCCTTTTCATCTTTGCA 520

RESULT 15
 US-09-774-639-99/c
 ; Sequence 99, Application US/09774639
 ; Publication No. US2003000355A1
 ; GENERAL INFORMATION:
 ; APPLICANT: Rosen et al.
 ; TITLE OF INVENTION: 90 Human Secreted Proteins
 ; FILE REFERENCE: P2013P1
 ; CURRENT APPLICATION NUMBER: US/09/774,639
 ; CURRENT FILING DATE: 2001-07-09
 ; PRIOR APPLICATION NUMBER: EARLIER APPLICATION NUMBER: 09/244,112
 ; PRIOR FILING DATE: EARLIER FILING DATE: 1999-02-04
 ; NUMBER OF SEQ ID NOS: 371
 ; SOFTWARE: Patentln Ver. 2.0
 ; SEQ ID NO 99
 ; LENGTH: 944
 ; TYPE: DNA
 ; ORGANISM: Homo sapiens
 ; FEATURE:
 ; NAME/KEY: SITE
 ; LOCATION: (13)
 ; OTHER INFORMATION: n equals a,t,g, or c
 ; NAME/KEY: SITE
 ; LOCATION: (486)
 ; OTHER INFORMATION: n equals a,t,g, or c
 ; NAME/KEY: SITE
 ; LOCATION: (934)
 ; OTHER INFORMATION: n equals a,t,g, or c
 US-09-774-639-99

Query Match 4.3%; Score 17; DB 9; Length 944;
 Best Local Similarity 100.0%; Pred. No. 29;
 Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 246 CTCACCTGCTTCCTTG 262
|||||
Db 515 CTCACCTGCTTCCTTG 499

Search completed: April 25, 2003, 02:12:39
Job time : 286.231 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - nucleic search, using frame_plus.p2n model

Run on: April 24, 2003, 22:30:38 ; Search time 1361 Seconds
(without alignments)
791.186 Million cell updates/sec

Title: US-09-513-999c-7869_COPY_1_37
Perfect score: 193
Sequence: 1 MGSEFALODSFSSLOGLLGPEYKVLGLGVCLSGCSCT 37

Scoring table:
BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

ched: 2054640 segs, 14551402878 residues
Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODE=frame+ p2n.model -DEV=xlh
-Q/cgn2.1/USPTO.spool/US09513999/unat_18042003.170936_28357/app-query.fasta.1.199
-DB=GenEmbl -QFMT=fastap -SUFFIX=p2n.rge -MINMATCH=0.1 -LOOPEXT=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=-1 -MATRIX=biosum62 -TRANS=human40.cdi -LIST=45
-DOCALIGN=200 -THR.SCORE=pct -THR.MAX=100 -THR.MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=fto -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09513999.ecgn.1.1.1687@unat.18042003.170936.28357 -NCPU=6 -ICPU=3
-NO_XUPXY -NO_MMAP -LARGEQUERY -NEG.SCORES=0 -WAIT -LONGLOC -DEV.TIMEOUT=120
-WARN.TIMEOUT=30 -THREDS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :
1: gb.ba:*
2: gb.htg:*
3: gb.in:*
4: gb.om:*
5: gb.ov:*
6: gb.pat:*
7: gb.ph:*
8: gb.pl:*
9: gb.pr:*
10: gb.ro:*
11: gb.sts:*
12: gb.sy:*
13: gb.un:*
14: gb.vi:*
15: em.ba:*
16: em.fun:*
17: em.hum:*
18: em.in:*
19: em.mu:*
20: em.om:*
21: em.or:*
22: em.ov:*
23: em.pat:*
24: em.ph:*
25: em.pl:*
26: em.ro:*
27: em.sts:*
28: em.un:*

29: em.vi:*
30: em.htg.hum:*
31: em.htg.inv:*
32: em.htg.other:*
33: em.htg.mus:*
34: em.htg.pin:*
35: em.htg.rod:*
36: em.htg.mam:*
37: em.htg.vrt:*
38: em.sy:*
39: em.htgo.hum:*
40: em.htgo.mus:*
41: em.htgo.other:*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
c 1	193	100.0	169620	2 AC012674	AC012674 Homo sapi
c 2	193	100.0	199289	2 AC012378	AC012378 Homo sapi
c 3	129	66.8	152313	9 AL591594	AL591594 Human DNA
c 4	129	66.8	163314	2 AL590557	AL590557 Homo sapi
c 5	127	65.8	143372	9 AL137847	AL137847 Human DNA
c 6	117	60.6	148290	9 AC107979	AC107979 Homo sapi
c 7	117	60.6	165649	9 AC103996	AC103996 Homo sapi
c 8	117	60.6	169861	2 AC126324	AC126324 Homo sapi
c 9	117	60.6	192826	2 AC090762	AC090762 Homo sapi
c 10	116	60.1	131347	2 AC002421	AC002421 Homo sapi
c 11	116	60.1	170623	2 AL391375	AL391375 Human DNA
c 12	114	59.1	68314	2 AC126345	AC126345 Homo sapi
c 13	114	59.1	143717	2 AC069245	AC069245 Homo sapi
c 14	114	59.1	152354	2 AC016472	AC016472 Homo sapi
c 15	114	59.1	160066	2 AC027295	AC027295 Homo sapi
c 16	114	59.1	163475	9 AC078815	AC078815 Homo sapi
c 17	114	59.1	200543	2 AC016726	AC016726 Homo sapi
c 18	113	58.5	57662	2 AC107969	AC107969 Homo sapi
c 19	113	58.5	98360	9 HSD247C2	AL049713 Human DNA
c 20	113	58.5	145264	9 AC107939	AC107939 Homo sapi
c 21	112	58.0	147820	9 AC092837	AC092837 Homo sapi
c 22	112	58.0	199038	2 AC116565	AC116565 Homo sapi
c 23	109	56.5	86314	9 AC109592	AC109592 Homo sapi
c 24	109	56.5	159475	2 AC021378	AC021378 Homo sapi
c 25	109	56.5	176689	9 AL162414	AL162414 Human DNA
c 26	108	56.0	134760	9 AC099484	AC099484 Homo sapi
c 27	108	56.0	166706	9 AC068875	AC068875 Homo sapi
c 28	108	56.0	207408	2 AC068618	AC068618 Homo sapi
c 29	108	56.0	207548	2 AC087283	AC087283 Homo sapi
c 30	107	55.4	103138	2 AC115094	AC115094 Homo sapi
c 31	107	55.4	143200	9 AC008413	AC008413 Homo sapi
c 32	107	55.4	231948	2 AC113346	AC113346 Homo sapi
c 33	106.5	55.2	146059	2 AC019030	AC019030 Homo sapi
c 34	106	54.9	177447	2 AC104687	AC104687 Homo sapi
c 35	106	54.9	236281	9 AC004673	AC004673 Homo sapi
c 36	105	54.4	80417	9 AL512659	AL512659 Human DNA
c 37	105	54.4	108248	9 HS326T13	AL022158 Homo sapi
c 38	105	54.4	162778	9 AC009559	AC009559 Homo sapi
c 39	104	53.9	24725	9 AL392090	AL392090 Human DNA
c 40	104	53.9	120418	2 AP002509	AP002509 Homo sapi
c 41	104	53.9	147288	9 AL355812	AL355812 Human DNA
c 42	104	53.9	163231	9 AL162575	AL162575 Human DNA
c 43	104	53.9	175832	9 AC023080	AC023080 Homo sapi
c 44	104	53.9	187847	2 AC023155	AC023155 Homo sapi
c 45	102	52.8	66489	2 AC100766	AC100766 Homo sapi

RESULT 1

ALIGNMENTS

AC012674/c
LOCUS AC012674 169620 bp DNA linear HTG 07-SEP-2000
DEFINITION Homo sapiens chromosome 3 clone RP1-458H3, WORKING DRAFT SEQUENCE.
AC012674
18 unordered pieces.
ACCESSION AC012674.10 GI:9719580
VERSION HTG: HTGS_PHA5E1; HTGS_DRAFT.
KEYWORDS Homo sapiens.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS Muzny, D.M., Adams, C., Bailey, M., Barbara, J., Blankenburg, K.,
Bodet, B., Bouck, J., Bowie, S., Brooks, A., Buhay, C., Bunac, C.,
Buckett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C.,
David, R., Delgado, O., Deshazo, D., Ding, Y., Domah-Rashid, N.,
Dugan-Rocha, S., Durbin, K.J., Fernandez, C., Frettaguto, D.,
Forcum-Tansey, J., Frantz, P., Ganes, R., Gorrell, J.H., Gorrell, L.L.,
Gouveira, W., Hosak, H., Jackson, L.E., Hodgson, A., Hogues, M.,
Holloway, C., Harris, K., Hernandez, J., Hodgson, A., Hognes, M.,
Kelly, S., Kondejewski, N., Kong, Y., Kovar, C., Leal, B., Li, Z.,
Lichter, O., Liu, J., Liu, W., Logan, O., Lozano, R.J., Lu, J.,
Lucier, R., Martin, R., Martinez, C., McLeod, M.P., Mei, G., Morgan, M.,
Morris, S., Nash, S., Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S.,
Oswal, G., Parish, B., Paxton, S., Payton, B., Perez, L., Pu, L.,
Quiles, M., Reiter, D., Rives, M., Samuel, S., Say, J., Scherer, S.,
Shah, E., Shen, H., Simon, M., Sparks, A., Stamps, A., Sugeng, S.,
Taber, P., Taylor, T., Vasquez, L., Vinson, R., Vo, O., Wabash, M.,
Wellington, S., Weinstein, G., Weinstein, I.R., Williamson, A.,
Worley, K., Wren, J., Wrenford, G., Yu, W., Zhou, X., Nelson, D., and
Gibbs, R.
Direct Submission
Unpublished
2 (bases 1 to 169620)
Worley, K.C.
Direct Submission
Submitted (03-NOV-1999) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Aug 7, 2000 this sequence version replaced gi:8705345.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HMOG
Center clone name: RP1-458H3
----- Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 139025 bases at least Q40
Consensus quality: 154842 bases at least Q30
Consensus quality: 159725 bases at least Q20
Estimated insert size: 162720; sum-of-contris estimation
Estimated insert size: 171608; agarose-1p estimation
Quality coverage: 3.9x in Q20 bases; agarose-1p estimation
Quality coverage: 4.1x in Q20 bases; sum-of-contris estimation

* NOTE: Estimated insert size may differ from sequence length
* (see <http://www.hgsc.bcm.tmc.edu/docs/genbank-draft-data.html>).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
1
28689: contig of 28689 bp in length
*
28690 28789: gap of unknown length
*
50832 50833: contig of 22043 bp in length
*
50833 50932: gap of unknown length
*
50933 69144: contig of 18212 bp in length

* 69145 69244: gap of unknown length
* 69245 84204: contig of 14660 bp in length
* 84205 84304: gap of unknown length
* 84305 94667: contig of 10463 bp in length
* 94668 94767: gap of unknown length
* 94768 107261: contig of 12494 bp in length
* 107262 107361: gap of unknown length
* 107362 117550: contig of 10189 bp in length
* 117551 117650: gap of unknown length
* 117651 126939: contig of 9289 bp in length
* 126940 127039: gap of unknown length
* 127040 135040: contig of 8001 bp in length
* 135041 135140: gap of unknown length
* 135141 141639: contig of 6499 bp in length
* 141640 141739: gap of unknown length
* 141740 149558: contig of 7819 bp in length
* 149559 149658: gap of unknown length
* 149659 154562: contig of 4904 bp in length
* 154563 154662: gap of unknown length
* 154663 158987: contig of 4325 bp in length
* 158988 159087: gap of unknown length
* 159088 162376: contig of 3289 bp in length
* 162377 162476: gap of unknown length
* 162477 165191: contig of 2715 bp in length
* 165192 165291: gap of unknown length
* 165292 167173: contig of 1882 bp in length
* 167174 167273: gap of unknown length
* 167274 168393: contig of 1120 bp in length
* 168394 168493: gap of unknown length
* 168494 169620: contig of 1127 bp in length.
Location/Qualifiers
1. 169620
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="3"
/clone="RP1-458H3"
BASE COUNT 52024 a 33180 c 32128 g 50322 t 1966 others
ORIGIN

Alignment Scores:
Pred. No.: 9,18e-18 Length: 169620
Score: 193.00 Matches: 37
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 2 Gaps: 0
US-09-513-999c-COPY_1_37 (1-37) x AC012674 (1-169620)
QY 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyPro 20
|||||
Db 87441 ATGGGCGAATCTTTTCCTTCGACGAGATCTTTTATCTTTTCGACGAGACTTCTGGGGCCG 87382
QY 21 GltTyrValIysLeuLeuGlyLeuGlyCysValCysLeuSerGlyCysSerThr 37
|||||
Db 87381 GAGTATGTAACCTCTGCTGTGTGTCTGTGACGAGGCTGCTACT 87331
RESULT 2
AC012378/c AC012378 199289 bp DNA linear PRI 09-AUG-2001
LOCUS AC012378 Homo sapiens chromosome 15 clone RP11-420M1 map 15q21.3, complete
DEFINITION sequence.
ACCESSION AC012378
VERSION AC012378.10 GI:15145648
KEYWORDS HTG.
SOURCE Homo sapiens.
ORGANISM Homo sapiens.
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE
AUTHORS Rowen, L., Madan, A., Qin, S., Baradarani, L., Birditt, B., Bloom, S.,
Burke, J., Dors, M., Fleetwood, P., Kaur, A., Madan, A., Nesbitt, R.,
Pate, D., and Hood, L.

TITLE Sequencing of human chromosome 15 D15S146-D15S117 region
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 199289)
AUTHORS Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B., Bloom,S., Dors,M., Dickhoff,R., Harrison,G., James,R., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T. and Hood,L.
JOURNAL Submitted (26-OCT-1999) Multimegabase Sequencing Center, University of Washington, PO Box 357730, Seattle, WA 98195, USA
REFERENCE 3 (bases 1 to 199289)
AUTHORS Rowen,L., Madan,A., Qin,S., Baradarani,L., Birditt,B., Bloom,S., Burke,J., Dors,M., Fleetwood,P., Kaur,A., Madan,A., Nesbitt,R., Pate,D. and Hood,L.
JOURNAL Submitted (09-AUG-2001) Multimegabase Sequencing Center, Institute for Systems Biology, 4225 Roosevelt Way NE, Suite 200, Seattle, WA 98103, USA
COMMENT On Aug 9, 2001 this sequence version replaced gi:13775292.
----- Genome Center
Center: Multimegabase Sequencing Center
Center code: UWMSC
Web site: http://chroma.mbt.washington.edu/msg_www
Contact: leetowensystemsbiology.org
----- Summary Statistics
Sequencing vector: pUC18; 108752
Chemistry: Dye-terminator Big Dye; 90% of reads
Chemistry: Dye-primer Big Dye; 10% of reads
Assembly program: Phrap; version 0.990399

NOTE: Data from overlapping clones AC022083 [Drafting center: UWMSC], AC012674 [Drafting center: BCM] and AC009997 [drafting center: UWMSC] were added for finishing.

FEATURES

source

1. 199289
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="15"
/map="15q21.3"
/clone="RP11-420M1"
/clone_lib="RPCT human bac library 11"
/note="This clone overlaps CTD-2137J4 AC022083, RP11-458I3 AC012674 and RP11-291H24 AC009997. Data from overlapping BACs were added and the consensus sequence determined from RP11-420M1 to the extent possible."
1. 165029
/note="Overlap with RP11-458H3, AC012674"
1. 11647
/note="Overlap with CTD-2137J4, AC022083"
49933. .50105
/note="Low quality data"
52173. .52233
/note="Low quality data"
52381. .52385
/note="Low quality data"
52412. .52483
/note="Low quality data"
52541. .52570
/note="Low quality data"
53242. .53257
/note="Low quality data"
53940. .54690
/note="Sequence data generated from subcloned PCR product"
72103. .72105
/note="Low quality data"
1503032. .155038
/note="Low quality data"
166787. .199289
/note="Overlap with RP11-291H24, AC009997"
BASE COUNT 63165 a 40338 c 36951 g 58835 t
ORIGIN

Alignment Scores:

Pred. No.: 1.08e-17 Length: 199289
Score: 193.00 Matches: 37
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
DB: 9 Gaps: 0
US-09-513-999c_Copy_1_37 (1-37) x AC012378 (1-199289)

QY 1 MetClyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyPro 20

Db 14756 ATGGTGGAATCTTTTGGCTTCAGGATTCCTTTTCATCTTTCAGGAGCTTCGGGCGG 14697

QY 21 GltTyrValIysLeuLeuGlyLeuCysValCysLeuSerGlyCysSerThr 37

Db 14696 GAGTATGTAACCTCCCTGGCTCTGTGTGTGCTAGTGGCTGCTTACT 14646

RESULT 3

AL591594/c 152313 bp DNA linear PRI 16-NOV-2001

LOCUS Human DNA sequence from clone RP11-424N15 on chromosome 1, complete

DEFINITION Sequence.

ACCESSION AL591594 GI:16944148

VERSION AL591594.9 HTG.

KEYWORDS HTG.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

TITLE 1 (bases 1 to 152313)

JOURNAL Direct Submission

Submitted (16-NOV-2001) Wellcome Trust Sanger Institute, Hinxton,

Cambridgeshire, CB10 1SA, UK. E-mail enquiries:

humquerry@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk

On Nov 15, 2001 this sequence version replaced gi:16605741.

During sequence assembly data is compared from overlapping clones.

Where differences are found these are annotated as variations

together with a note of the overlapping clone name. Note that the

variation annotation may not be found in the sequence submission

corresponding to the overlapping clone, as we submit sequences with

only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all

regions were either double-stranded or sequenced with an alternate

chemistry or covered by high quality data (i.e., phred quality >=

30); an attempt was made to resolve all sequencing problems, such

as compressions and repeats; all regions were covered by at least

one plasmid subclone or more than one M13 subclone; and the

assembly was confirmed by restriction digest. The following

abbreviations are used to associate primary accession numbers given

in the feature table with their source databases: Em: EMBL; Sw: SWISSPROT; Tr: TrEMBL; Wp: WORMPEP; Information on the WORMPEP

database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence

was generated from part of bacterial clone contigs of human

chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping

Group. Further information can be found at

http://www.sanger.ac.uk/HGP/Chrl

RP11-424N15 is from the library RPCT-11.2 constructed by the group

of Pieter de Jong. For further details see

http://www.choi1.org/bacpac/home.htm

VECTOR: PBACe3.6

This sequence is the entire insert of clone RP11-424N15 The true

right end of clone RP11-518J10 is at 64264 in this sequence.

FEATURES

source

1. 152313
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="1"
/clone="RP11-424N15"
/clone_lib="RPCT-11.2"
/note="Sequence from overlapping clone RP11-24C8
18821. .18877

```

BASE COUNT      45419 a 29865 c 29040 g 47989 t          (AL590557). Assembly confirmed by restriction digest."
ORIGIN

Alignment Scores:
Pred. No.:           5.03e-08              Length:             152313
Score:               129.00                Matches:            23
Percent Similarity:  92.86%                 Conservative:       3
Best local Similarity: 82.14%                Mismatches:        2
Query Match:         66.84%                  Indels:            0
DB:                  9                      Gaps:              0

US-09-513-999C-7869_COPY_1_37 (1-37) x AL591594 (1-152313)
OY      9   AspSerPheSerSerLeuGlnGlyLeuLeuGlyProGlnTyValIysLeuLeuGlyLeu 28
||| |||||:::||| ::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||::|||
Db 100967  GACAAATTTTCACCTTGCCAGGGCGTCTGTGGTCCACGTATGTAAACTCTGGGACC TC 100908

OY      29  CysValCysLeuSerCylCysSer 36
|||||TTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTTT
Db 100907  TGCCTGTGCCCTGAGTGGCTGCTCT 100884

RESULT 4
AL590557 LOCUS Homo sapiens chromosome 1 clone RP11-24C8, *** SEQUENCING IN PROGRESS ***, 8 unordered pieces.
AL590557 ACCESSION AL590557 GI:13992136
VERSION HTG; HTGS_PHASE1; HTGS_CANCELLED.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 163314)
McJarry, K.
Direct Submission
Submitted (20-JUL-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
On May 8, 2001 this sequence version replaced gi:13990622.
----- Genome Center -----
Center: Sanger Centre
Center code: SC
Web site: http://www.sanger.ac.uk
Contact: humquery@sanger.ac.uk
----- Project Information -----
Center project name: ba24c8
----- Summary Statistics -----
Sequencing program: XGAP4: version 4.5
Chemistry: Dye-terminator Big Dye; 100% of reads
Consensus quality: 161985 bases at least Q40
Consensus quality: 161940 bases at least Q30
Consensus quality: 162264 bases at least Q20
Insert size: 162614; sum-of-contigs
Insert size: 164357; 4.9% error; agarose-fp
Quality coverage: 6.63x in Q20 bases; sum-of-contigs Quality
coverage: 6.67x in Q20 bases; agarose-fp
----- NOTE: This is a 'working draft' sequence. It currently consists of 8 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as * runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.
*****
1 16091: contig of 16091 bp in length
* 16092 16191: gap of 100 bp
* 16192 51184: contig of 34993 bp in length
* 51185 51284: gap of 100 bp
* 51285 67941: contig of 16657 bp in length

```

```

67942 68041: gap of 100 bp
*
* 68042 103921: contig of 35880 bp in length
*
* 103922 104021: gap of 100 bp
*
* 104022 107020: contig of 2999 bp in length
*
* 107021 107120: gap of 100 bp
*
* 107121 123748: contig of 16628 bp in length
*
* 123749 123848: gap of 100 bp
*
* 123849 137091: contig of 13243 bp in length
*
* 137092 137191: gap of 100 bp
*
* 137192 163314: contig of 26123 bp in length.
Location/Qualifiers
source
1..163314
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="1"
/clone="RP11-24C8"
/clone_lib="RPC1-11.1"
1..16091
misc_feature
/note="assembly_fragment:00603
fragment_chain:1"
16192..51184
misc_feature
/note="assembly_fragment:01328
fragment_chain:1"
51285..67941
misc_feature
/note="assembly_fragment:01579
fragment_chain:1"
68042..103921
misc_feature
/note="assembly_fragment:00704
fragment_chain:1"
104022..107020
misc_feature
/note="assembly_fragment:02225
fragment_chain:1"
107121..123748
misc_feature
/note="assembly_fragment:02172
fragment_chain:1"
123849..137091
misc_feature
/note="assembly_fragment:01998
fragment_chain:1"
137192..163314
misc_feature
/note="assembly_fragment:03121"
BASE COUNT 51215 a 31325 c 32024 g 48050 t 700 others
ORIGIN
Alignment Scores:
Pred. No.: 5,39e+08 Length: 163314
Score: 129..00 Matches: 23
Percent Similarity: 92.86% Conservative: 3
Best Local Similarity: 82.14% Mismatches: 2
Query Match: 66.84% Indels: 0
DB: 2 Gaps: 0
US-09-513-999C-7869_COPY_1_37 (1-37) x ALS90557 (1-163314)
QY 9 AspSerPheSerLeuGlnGlyLeuEngLyProGluTyValIysLeuEngLyLeu 28
||| | | | |
||||
|||||
||||||
|||||||
|||||||
|||||||
|||||||
|||||||
|||||||
|||||||
Db 55961 GACCAATTTTCCACCCTTGCGGTCCGTCGCACAGTAATAAACTCCTGGGCCCTC 56020
|||||
|||||
|||||
|||||
Oy 29 CysValGlySerGlyCysSer 36
|||||
|||||
|||||
|||||
Db 56021 TGC GTGCTGAGTG GCTGCTCT 56044

RESULT 5
AL137847/c 143372 bp DNA linear PRI 16-NOV-2001
LOCUS AL137847
DEFINITION Human DNA sequence from clone RP11-439K3 on chromosome 9q22.2-31.1.1.
complete sequence.
ACCESSION AL137847
VERSION AL137847.12 GI:16973786
KEYWORDS HTG.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
```


ACCESSION	ACI03996
VERSION	ACT03996.7
KEYWORDS	GI:21637504
SOURCE	Hrg.
ORGANISM	human.
REFERENCE	Homo sapiens
AUTHORS	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 165649)
JOURNAL	Bliren,B., Nusbaum,C. and Lander,E. Homo sapiens chromosome 15, clone RP11-76E17 Unpublished 2 (bases 1 to 165649)
REFERENCE	Bliren,B., Linton,L., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Boukhalter,B., Brown,M., Camarata,J., Campolongo,A., Chang,J., Chazarro,B., Choepe,Y., Collangelo,M., Collins,S., Collamore,A., Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N., Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Larocque,K., Lamazeres,R., Landers,T., Lehoczy,J., Levine,R., Liu,G., Maclean,C., Macdonald,P., Major,J., Margulis,N., Matthews,C., McCarthy,M., McKernan,K., McPeeters,R., Meldrum,J., Menues,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schnuppack,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N., Strassman,K., Subramanian,A., Talamas,J., Testaye,S., Theodore,J., Topham,K., Travers,M., Travis,N., Trillio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and zody,M.
TITLE	Direct Submission
JOURNAL	Submitted (01-DEC-2001) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA 3 (bases 1 to 165649)
REFERENCE	Bliren,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhalter,B., Camarata,J., Chang,J., Chazarro,B., Choepe,Y., Collamore,A., Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C., McCarthy,M., Meldrum,J., Menues,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Roy,A., Schauer,S., Schnuppack,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J., Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and zody,M.
TITLE	Direct Submission
JOURNAL	Submitted (26-JUN-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA 4 (bases 1 to 165649)
REFERENCE	Bliren,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhalter,B., Camarata,J., Chang,J., Chazarro,B., Choepe,Y., Collamore,A., Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J., Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K., Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C., McCarthy,M., Meldrum,J., Menues,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P., Roman,J., Roy,A., Schauer,S., Schnuppack,R., Seaman,S., Severy,P., Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J., Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and zody,M.

TITLE	Roman, J., Roy, A., Schauer, S., Schuppach, R., Seaman, S., Severy, P., Smith, C., Spencer, B., Strange, R., Thomann, N., Stojanovic, N., Talamas, J., Testa, E., Thedore, J., Topham, K., Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
JOURNAL	Direct Submission
COMMENT	Submitted (01-JUL-2002) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA On Jul 1, 2002 this sequence version replaced g1:21592191. All repeats were identified using RepeatMasker: Smit, A.F.A. & Green, P. (1996-1997) http://ftp.genome.washington.edu/RM/RepeatMasker.html
FEATURES	----- Genome Center Center: Whitehead Institute/ MIT Center for Genome Research Center code: WIBR Web site: http://www-seq.wi.mit.edu Contact: sequence_submissions@genome.wi.mit.edu ----- Project Information Center project name: L21917 Center clone name: 76_E_17 -----
source	Location/Qualifiers 1..165649 /organism="Homo sapiens" /db_xref="taxon:9606" /chromosome="15" /map="15" /clone="RP11-76E17" /clone_1b="RPC1-11 Human Male BAC" /complement(2..865) /rpt_family="L1PA13" /complement(864..894) /rpt_family="L1PA13" /complement(895..1248) /rpt_family="THE1A" /complement(1249..1504) /rpt_family="L1PA13" 1505..1626 /rpt_family="A1uSx" 1627..1659 /rpt_family="A1uSx" 1660..1831 /rpt_family="A1uSx" /complement(1832..3332) /rpt_family="L1PA13" /complement(3340..3699) /rpt_family="L1uMC" /complement(4461..4755) /rpt_family="A1uSx" 5027..5397 /rpt_family="L2" /complement(5650..6330) /rpt_family="L1uME1" /complement(6343..6460) /rpt_family="L1uME1" 7214..7319 /rpt_family="L1uMC3" 7354..7655 /rpt_family="L1uMC3" 7659..7743 /rpt_family="L1PA10" /complement(7745..7843) /rpt_family="A1uSp/q" 7844..14156 /rpt_family="L1PA10" 14198..14328 /rpt_family="A1uY" 14340..14503 /rpt_family="(TA)n" 14507..15032 /rpt_family="L1uMC3" 15036..15220 /rpt_family="L1uMC3"

```

repeat_region complement(15218..16449)
repeat_region /rpt_family="L1PA4"
16450..18154
/rpt_family="L1PA4"
repeat_region 18155..19210
/rpt_family="L1MC3"
repeat_region 19211..19266
/rpt_family="(TA)n"
repeat_region 19267..19372
/rpt_family="L1MC3"
repeat_region 19401..19476
/rpt_family="(TTATA)n"
repeat_region 19483..19546
/rpt_family="(CAATATA)n"
repeat_region 19596..19625
/rpt_family="AT-rich"
repeat_region 19696..19750
/rpt_family="GA-rich"
repeat_region 19752..19920
/rpt_family="L1MD3"
repeat_region 19994..20102
/rpt_family="L2"
20049..20114
/ncore="single clone coverage"
repeat_region 20485..20655
/rpt_family="MIR3"
repeat_region complement(21285..21441)
/rpt_family="MIR"
21496..21717
/rpt_family="L2"
repeat_region complement(21720..21796)
/rpt_family="MIR"
repeat_region complement(21943..22131)
/rpt_family="MIR"
repeat_region 23082..23195
/rpt_family="L2"
23198..23248
/rpt_family="GA-rich"
repeat_region 23267..23333
/rpt_family="(CAAT)n"
repeat_region complement(24465..24833)
/rpt_family="M1T1A2"
26142..26334
/rpt_family="MIR"
repeat_region complement(27659..27811)
/rpt_family="MIR"
repeat_region complement(28811..28880)
/rpt_family="MIR"
repeat_region complement(29026..29269)
/rpt_family="MIR"
repeat_region complement(29673..29706)
/rpt_family="MSTB"

Alignment Scores:
Pred. No.: 3.74e-06 Length: 165649
Score: 117.00 Matches: 21
Percent Similarity: 85.71% Conservative: 4
Best Local Similarity: 75.00% Mismatches: 4
Query Match: 60.62% Indels: 0
DB: 9 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AC103996 (1-165649)
OY 9 AspserPheserSeuLeuGngLyLeuGngLyProGluTyValYsLeuLeuGngLyLeu 28
DB 8153 GATGATCTTCCACCTCACTGCGGATCCTGGGCGCAGATATATAAACTCTCGGTCCTC 8094
OY 29 CysValCysLeuSerGlyCysSer 36
DB 8093 TGTGTGCTGCTGAGTGGCTGCTCA 8070
RESULT 8
AC126324/c

```

```

LOCUS AC126324 169861 bp DNA linear HTG 06-AUG-2002
DEFINITION Homo sapiens chromosome 11 clone RP11-358H20 map 11. *** SEQUENCING
IN PROGRESS ***; 2 ordered pieces.
ACCESSION AC126324
VERSION AC126324.2 GI:22123018
KEYWORDS HTG; HTGS_PHASE2; HTGS_FULFILLTOP; HTGS_ACTIVEFIN.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 169861)
Birren,B., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 11, clone RP11-358H20
Unpublished
2 (bases 1 to 169861)
Birren,B., Nusbaum,C., Lander,E., All,A., Allen,N., Anderson,S.,
Barra,N., Bastien,V., Bloom,T., Boguslavsky,L., Bookhalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., Deatrellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-roh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,D., Matthews,C.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
Phunkhang,P., Pierre,N., Raymond,S., Schnpbach,R., Seaman,S., Severy,P.,
Smit,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Willson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (05-JUL-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 169861)
Birren,B., Bastien,V., Bloom,T., Boguslavsky,L., Bookhalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., Deatrellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-roh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,D., Matthews,C.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
Phunkhang,P., Pierre,N., Raymond,S., Schnpbach,R., Seaman,S., Severy,P.,
Smit,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Willson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (06-AUG-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Aug 6, 2002 this sequence version replaced g1:21699255.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RN/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L27749
Center clone name: 358_H_20

```

* NOTE: This is a 'working draft' sequence. It currently consists of 2 contigs. Gaps between the contigs

* are represented as runs of N. The order of the pieces
 * is believed to be correct as given, however the sizes
 * of the gaps between them are based on estimates that have
 * provided by the submitter.
 * This sequence will be replaced
 * by the finished sequence as soon as it is available and
 * the accession number will be preserved.
 * 1 71322: contig of 71322 bp in length
 * 71323 71422: gap of 100 bp
 * 71423 169861: contig of 98439 bp in length.

FEATURES

source

1..169861
 /organism="Homo sapiens"
 /db_xref="taxon:9606"
 /chromosome="11"
 /map="11"
 /clone="RP11-358H20"
 /clone_lib="RPCT-11 Human Male BAC"

BASE COUNT 55771 a 33379 c 31394 g 49217 t 100 others

ORIGIN

Alignment Scores:

pred. No.: 3.83e-06 Length: 169861
 Score: 117.00 Matches: 23
 Percent Similarity: 79.41% Conservatave: 4
 Best Local Similarity: 67.65% Mismatches: 7
 Query Match: 60.62% Indels: 0
 DB: 2 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AC126324 (1-169861)

QY 3 GlycerPhlealLeuGlnAspSerPheSerLeuGlnGlyLeuGlyProGluTyr 22

Db 73373 GGACCTTCACGCCCTGCCTGAGTTCAGACAGCACTGTGTTGGCTTCCTGGCCACAGCTTT 73314

QY 23 ValTysLeuLeuGlyLeuGlyCysValcysLeuSerGlyCysSer 36

Db 73313 GTPAAGCTCTGCTGCTGTGTGTGTGACGACACTGCTCT 73272

RESULT 9

AC090762/c

LOCUS AC090762 192826 bp DNA linear PRI 28-FEB-2002
 DEFINITION Homo sapiens chromosome 15, clone RP11-387B8, complete sequence.

AC090762

AC090762.9 GI:18997378

KEYWORDS

SOURCE

ORGANISM

Homo sapiens.
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

1 (bases 1 to 192826)

Birren,B., Linton,L., Nusbaum,C. and Lander,E.

Homo sapiens chromosome 15, clone RP11-387B8

Unpublished

2 (bases 1 to 192826)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,

Barna,N., Bastien,V., Boguslavsky,L., Bouckhalter,B., Brown,A.,

Camata,J., Campopiano,A., Choepel,Y., Colangelo,M., Collins,S.,

Collimore,A., Cooke,P., Dearlano,K., Dewar,K., Diaz,J.S.,

Dodge,S., Fato,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J.,

Gardyna,S., Ginde,S., Goyette,M., Graham,L., Grand-Pierre,N.,

Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,

Jones,C., Karatas,A., Lacroque,K., Lamazares,R., Landers,T.,

Lehoczy,J., Levine,R., Liu,G., Maclean,C., Macdonald,P.,

Marguis,N., Matthews,C., McCarthy,M., McKernan,P., McKernan,K.,

McPheeers,R., Meldrim,J., Menus,L., Mihova,T., Mlenga,V.,

Murphy,T., Naylor,J., Nguyen,C., Norbu,C., Norman,C.H.,

O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,

Phunkhang,P., Pierre,N., Pollara,V., Raymond,C., Retta,R.,

Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J., Roselli,M.,

Roy,A., Santos,R., Schauer,S., Schupack,R., Seaman,S., Severy,P.,

Sougne,N., Spencer,B., Stange-Thomann,N., Stojanovic,N.,

Strauss,N., Subramanian,A., Talamas,J., Testaye,S., Theodore,J.,

Travers,M., Travis,N., Trigilio,J., Vassiliev,H., Viel,R., Vo,A.,
 Wilson,B., Wu,X., Wyman,D., Ye,M.J., Young,G., Zainoun,J.,
 Zembek,L., Zimmer,A. and Zody,M.

TITLE

JOURNAL

REFERENCE

AUTHORS

Submitted (10-MAR-2001) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 3 (bases 1 to 192826)

Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
 Barna,N., Bastien,V., Boguslavsky,L., Bouckhalter,B., Brown,A.,
 Camarata,J., Campopiano,A., Chang,J., Chazaro,B., Choepel,Y.,
 Colangelo,M., Collins,S., Collimore,A., Cooke,A., Cooke,P.,
 Dearlano,K., Dewar,K., Diaz,J.S., Dodge,S., Fato,S., Ferreira,P.,
 Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Ginde,S., Goyette,M.,
 Graham,L., Grand-Pierre,N., Hagos,B., Horton,L., Hulme,W.,
 Iliev,I., Johnson,R., Jones,C., Kamat,A., Karatas,A., Kells,C.,
 Lacroque,K., Lamazares,R., Landers,T., Lehoczy,J., Levine,R.,
 Liu,G., Maclean,C., Macdonald,P., Major,J., Marguis,N.,
 Matthews,C., McCarthy,M., McKernan,P., McKernan,K., Meldrim,J.,
 Menus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,
 Nicol,R., Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P.,
 O'Neill,D., Oliver,J., Peterson,K., Phunkhang,P., Pierre,N.,
 Pollara,V., Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C.,
 Rogov,P., Roman,J., Roselli,M., Roy,A., Santos,R., Schauer,S.,
 Schupack,R., Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N.,
 Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Testaye,S.,
 Theodore,J., Topham,K., Travers,M., Travis,N., Trigilio,J.,
 Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,M.J.,
 Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.

Direct Submission
 Submitted (28-FEB-2002) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA

COMMENT

On Feb 28, 2002 this sequence version replaced gi:18377189.
 All repeats were identified using RepeatMasker:

Smith, A.F.A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research

Center: WtIBR

Web site: http://www-seq.wi.mit.edu

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information

Center project name: L12392

Center clone name: 387_E8

location/Qualifiers

1..192826

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="15"

/map="15"

/clone="RP11-387B8"

/clone_lib="RPCT-11 Human Male BAC"

/complement(864..1222)

/rpt_family="L1PB3"

1223..1252

/rpt_family="(TAG)n"

complement(1253..1492)

/rpt_family="L1PB3"

4287..4576

/rpt_family="AluB"

4671..5034

/rpt_family="THELB"

5130..5438

/rpt_family="AluSx"

5517..5662

/rpt_family="MIR"

5798..5827

/rpt_family="AT_rich"

5985..6114

/rpt_family="AluB"

6227..6253

/rpt_family="(CA)n"

```

repeat_region      /rpl_family="CT-rich"
                   34655..34960
/rpl_family="AluY"
repeat_region      complement(35296..35389)
                   /rpl_family="MERSB"
                   complement(35419..35517)
repeat_region      /rpl_family="L1MC4"
                   complement(35518..35823)
repeat_region      /rpl_family="AluSx"
                   complement(35824..36030)
repeat_region      /rpl_family="L1MC4"
                   36682..36758
repeat_region      /rpl_family="MERCA"
                   36975..37160
repeat_region      /rpl_family="MERSA"
                   38140..38353
repeat_region      /rpl_family="MERS6B"
                   38653..38656
repeat_region      /rpl_family="(TTTA)n"
repeat_region      complement(38657..40056)
repeat_region      /rpl_family="L1PA5"
repeat_region      complement(40060..40250)

Alignment Scores:
Pred. NO.:         4.34e-06           Length:        192826
Score:            117.00              Matches:        21
Percent Similarity: 85.71%             Conservative:   3
Best Local Similarity: 75.00%          Mismatches:    4
Query Match:       60.62%              Indels:        0
Gap:               9                  Gaps:          0

US-09-513-999C-7869_COPY_1_37 (1-37) x AC090762 (1-192826)

Qy     9 AspserrPheSerSerLeuGInglyLeuLeuGlyProGUtyrValylSLseuLeuGlyLeu 28
      ||| |:::| |:::| |:::| |:::| |:::| |:::| |:::| |:::| |:::| |:::|
Db 190989 GATGGATCTTCACCTCCACTCGGGGATCTCGGCCAGCATATAAAACCTCGGTCTC 190930

Qy     29 CysvalCysleuSerGlyCysSer 36
      ||||| ||||||| ||||||| ||||||| ||||||| ||||||| ||||||| |||||||
Db 190929 TGTGTGTGCTTGAGTGGCTGCTCA 190906


RESULT 10
LOCUS       AC002421                131347 bp      DNA      Linear      HTG 13-JUN-2002
DEFINITION Homo sapiens chromosome X clone pxMD1, *** SEQUENCING IN PROGRESS
ACCSSION   AC002421
VERSION    AC002421.2 GI:21405641
KEYWORDS   HTG; HTGS_PHASE1.
SOURCE     human.
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
Chen,E., Brownstein,B.H., States,D.J., Schlessinger,D. and
Mazzarella,R.
Direct Submission
Unpublished (1997)
2 (bases 1 to 131347)
Brownstein,B.H., States,D.J. and Mazzarella,R.
Direct Submission
Submitted (12-MUG-1997) Center for Genetics in Medicine, Box 8232,,
Washington University School of Medicine, 4566 Scott Avenue, St.
Louis, MO 63110, USA
On Jun 13, 2002 this sequence version replaced g1:2323248.
Current status of this project is available at:
'http://genome.wustl.edu/cgm/seq_projects.html'
Submitted by:
Elison Chen.
Advanced Center for Genetic Technology,
Applied Biosystems Division of Perlin Elmer Corp.,
850 Lincoln Center Drive,
Foster City, CA 94404 USA
```

	repeat_region	/rpt_family="CT-rich"	
		34655..34960	
	rpt_family="AluY"		
	complement(35296..35389)		
	repeat_region	/rpt_family="MER5B"	
		complement(35419..35517)	
	repeat_region	/rpt_family="LIMC4"	
		complement(35518..35823)	
	repeat_region	/rpt_family="ALUSX"	
		complement(35824..36030)	
	repeat_region	/rpt_family="LIMC4"	
		36682..36758	
	repeat_region	/rpt_family="MERSA"	
		36975..37160	
	repeat_region	/rpt_family="MERSA"	
		38140..38353	
	repeat_region	/rpt_family="MERS6B"	
		38653..38656	
	repeat_region	/rpt_family="(TTTA)n"	
		complement(38657..40056)	
	repeat_region	/rpt_family="LIPAS"	
		complement(40060..40250)	
Alignment Scores:			
Pred. NO.:	4.34e-06	Length:	192826
Score:	117.00	Matches:	21
Percent Similarity:	85.71%	Conservative:	3
Best Local Similarity:	75.00%	Mismatches:	4
Query Match:	60.62%	Indels:	0
Gaps:	9	Gaps:	0
 US-09-513-999C-7869_COPY_1_37 (1-37) x AC090762 (1-192826)			
Oy	9 AspsErPheSerSerLeuglnclyleuleuclyprogtutryValylSlsleulglyleu	28	
Db	190989 GATGGATCTTCACCCACTGCGGAGTCTTGCGGCCAGCATATAAAACCTCGGTCTC	190930	
Oy	29 CysvalcysleuserglyCysser	36	
Db	190929 TGTGTGTGCTTGAGTGCGTCTCA	190906	
RESULT 10			
LOCUS	AC002421	131347 bp	DNA Linear HTG 13-JUN-2002
DEFINITION	Homo sapiens chromosome X clone pxMD1,	*** SEQUENCING IN PROGRESS ***	
ACCSSION	AC002421		
VERSION	AC002421.2	GI:21405641	
KEYWORDS	HTG; HTOGS_PHASEI.		
SOURCE	human.		
ORGANISM	Homo sapiens		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;		
AUTHORS	Mammalia; Euarchonta; Primates; Catarrhini; Homnidae; Homo.		
TITLE	JOURNAL		
AUTHORS	Chen,E., Brownstein,B.H., States,D.J., Schlessinger,D. and Mazzarella,R.		
TITLE	Direct Submission		
JOURNAL	Submitted (12-MGC-1997) Center for Genetics in Medicine, Box 8232, Washington University School of Medicine, 4566 Scott Avenue, St. Louis, MO 63110, USA On Jun 13, 2002 this sequence version replaced g1:2323248. Current status of this project is available at: 'http://genome.wustl.edu/cgm/seq_projects.html' Submitted by: Elison Chen, Advanced Center for Genetic Technology, Applied Biosystems Division of Perlin Elmer Corp., 850 Lincoln Center Drive, Foster City, CA 94404 USA		
COMMENT			

e-mail: ellson@genseq.apldbio.com
and

Buddy Brownstein,
Center for Genetics in Medicine,
Washington University School of Medicine, Box 8232
4566 Scott Avenue,
St. Louis, MO 63110, USA
e-mail: buddy@genetics.wustl.edu
and

David J. States,
Institute for Biomedical Computing
Washington University in St. Louis
700 South Euclid Ave.
St. Louis, MO 63108 USA

e-mail: states@dbc.wustl.edu.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 4 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

* 1 22595: contig of 22595 bp in length
* 22596 22695: gap of 100 bp
* 22696 66983: contig of 44288 bp in length
* 66984 67083: gap of 100 bp
* 67084 122847: contig of 55764 bp in length
* 122848 122947: gap of 100 bp
* 122948 131347: contig of 8400 bp in length.

FEATURES

source

1. .131347
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/clone="PMXD1"

BASE COUNT 40590 a 23040 c 23923 g 43494 t 300 others
ORIGIN

Alignment Scores:

Pred. No.: 4.22e-06 Length: 131347
Score: 116.00 Matches: 21
Percent Similarity: 86.21% Conservative: 4
Best Local Similarity: 72.41% Mismatches: 4
Query Match: 60.10% Indels: 0
Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AC002421 (1-131347)

QY 9 AspSerPheSerSerLeuGlnGlyLeuLeuGlyProGluTyrValIysLeuGlyLeu 28

Db 103911 GATGGATCTGCCACCTTCCTGGGAATCTGGGCCAGAGTAAACTCTCGGCTCTC 103970

QY 29 CysValCysLeuSerGlyCysSerThr 37

Db 103971 TGTGTGTGCTGAGCAGCTGCTTCT 103997

RESULT 11

AL391375

LOCUS AL391375 170623 bp DNA linear PRI 02-FEB-2001
DEFINITION Human DNA sequence from clone RP11-375A20 on chromosome X, complete
sequence.

ACCESSION AL391375 GI:12657182

VERSION

KEYWORDS

HTG.

SOURCE

ORGANISM

Homo sapiens
human.
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.

REFERENCE 1 (bases 1 to 170623)

AUTHORS

Chapman, J.

Direct Submission

JOURNAL

Submitted (02-FEB-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk

COMMENT

On Feb 2, 2001 this sequence version replaced gi:11414631.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence has been finished according to sequence map criteria
as follows. An attempt is made to resolve all sequencing problems,
such as compressions and repeats, but not necessarily within known
annotated repeat sequence elements. Where the sequence is
ambiguous, there is an annotation using the 'unsure' feature key.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/projects/C.elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome X, constructed by the Sanger Centre Chromosome X Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/ChrX
RP11-375A20 is from the library RPCI-11.2 constructed by the group
of Pletier de Jong. For further details see
http://www.chori.org/bacpac/home.htm
VECTOR: pBACE3.6
This sequence is the entire insert of clone RP11-375A20.

FEATURES

source

1. .170623
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="X"
/clone="RP11-375A20"
/clone_11b="RPCI-11.2"
71147..71285
/note="Single clone region. Assembly confirmed by
restriction digest data."

BASE COUNT 53145 a 29784 c 30919 g 56775 t
ORIGIN

Alignment Scores:

Pred. No.: 5.47e-06 Length: 170623
Score: 116.00 Matches: 21
Percent Similarity: 86.21% Conservative: 4
Best Local Similarity: 72.41% Mismatches: 4
Query Match: 60.10% Indels: 0
Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AL391375 (1-170623)

QY 9 AspSerPheSerSerLeuGlnGlyLeuLeuGlyProGluTyrValIysLeuGlyLeu 28

Db 99926 GATGGATCTGCCACCTTCCTGGGAATCTGGGCCAGAGTAAACTCTCGGCTCTC 99985

QY 29 CysValCysLeuSerGlyCysSerThr 37

Db 99986 TGTGTGTGCTGAGCAGCTGCTTCT 100012

RESULT 12

AC126345/c

LOCUS AC126345 68314 bp DNA linear HTG 30-JUL-2002
DEFINITION Homo sapiens chromosome 11 clone RP11-100E23 map 11, LOW-PASS
SEQUENCE SAMPLING.

ACCESSION

VERSION

KEYWORDS

HTG: HTGS_PHASED.

SOURCE

ORGANISM

Homo sapiens
human.

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 68314)
Birren,B., Nusbaum,C. and Lander,E.
Homo sapiens chromosome 11, clone RP11-100E23
Unpublished
2 (bases 1 to 68314)
Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barra,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhalter,B.,
Camrata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faroo,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardina,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kanat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrum,J., Menus,L., Mihova,T., Miengs,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,B., Schupback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (05-JUL-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 68314)
Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barra,N., Bastien,V., Bloom,T., Boguslavskiy,L., Boukhalter,B.,
Camrata,J., Chang,J., Chazaro,B., Choepel,Y., Collymore,A.,
Cook,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faroo,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardina,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kanat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., Maclean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrum,J., Menus,L., Mihova,T., Miengs,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neill,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,B., Schupback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Testaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (30-JUL-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jul 30, 2002 this sequence version replaced gi:21699290.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
Project Information
Center project name: L27743
Center clone name: 100_E_23
NOTE: This record contains 84 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will

* be preserved. 728: contig of 728 bp in length
* 1 729 828: gap of 100 bp
* 829 1544: contig of 716 bp in length
* 1545 1644: gap of 100 bp
* 1645 2360: contig of 716 bp in length
* 2361 2460: gap of 100 bp
* 2461 3152: contig of 692 bp in length
* 3153 3252: gap of 100 bp
* 3253 3960: contig of 708 bp in length
* 3961 4060: gap of 100 bp
* 4061 4774: contig of 714 bp in length
* 4775 4874: gap of 100 bp
* 4875 5583: contig of 709 bp in length
* 5584 5683: gap of 100 bp
* 5684 6389: contig of 706 bp in length
* 6390 6489: gap of 100 bp
* 6490 7213: contig of 724 bp in length
* 7214 7313: gap of 100 bp
* 7314 8019: contig of 706 bp in length
* 8020 8119: gap of 100 bp
* 8120 8851: contig of 732 bp in length
* 8852 8951: gap of 100 bp
* 8952 9622: contig of 671 bp in length
* 9623 9722: gap of 100 bp
* 9723 10434: contig of 712 bp in length
* 10435 10534: gap of 100 bp
* 10535 11257: contig of 723 bp in length
* 11258 11357: gap of 100 bp
* 11358 12070: contig of 713 bp in length
* 12071 12170: gap of 100 bp
* 12171 12895: contig of 725 bp in length
* 12896 12995: gap of 100 bp
* 12996 13714: contig of 719 bp in length
* 13715 13814: gap of 100 bp
* 13815 14524: contig of 710 bp in length
* 14525 14624: gap of 100 bp
* 14625 15335: contig of 711 bp in length
* 15336 15435: gap of 100 bp
* 15436 16165: contig of 730 bp in length
* 16166 16265: gap of 100 bp
* 16266 16986: contig of 721 bp in length
* 16987 17086: gap of 100 bp
* 17087 17817: contig of 731 bp in length
* 17818 17917: gap of 100 bp
* 17918 18633: contig of 716 bp in length
* 18634 18733: gap of 100 bp
* 18734 19459: contig of 726 bp in length
* 19460 19559: gap of 100 bp
* 19560 20266: contig of 707 bp in length
* 20267 20366: gap of 100 bp
* 20367 21090: contig of 724 bp in length
* 21091 21190: gap of 100 bp
* 21191 21901: contig of 711 bp in length
* 21902 22001: gap of 100 bp
* 22002 22721: contig of 720 bp in length
* 22722 22821: gap of 100 bp
* 22822 23530: contig of 709 bp in length
* 23531 23630: gap of 100 bp
* 23631 24336: contig of 706 bp in length
* 24337 24436: gap of 100 bp
* 24437 25167: contig of 731 bp in length
* 25168 25267: gap of 100 bp
* 25268 25989: contig of 722 bp in length
* 25990 26089: gap of 100 bp
* 26090 26816: contig of 727 bp in length
* 26817 26916: gap of 100 bp
* 26917 27618: contig of 702 bp in length
* 27619 27718: gap of 100 bp
* 27719 28439: contig of 721 bp in length
* 28440 28539: gap of 100 bp
* 28540 29257: contig of 718 bp in length
* 29258 29357: gap of 100 bp

```
* 29358 30053: contig of 696 bp in length
* 30054 30153: gap of 100 bp
* 30154 30856: contig of 703 bp in length
* 30857 30956: gap of 100 bp
* 30957 31680: contig of 724 bp in length
* 31681 31780: gap of 100 bp
* 31781 32509: contig of 729 bp in length
* 32510 32609: gap of 100 bp
* 32610 33331: contig of 722 bp in length
* 33332 33431: gap of 100 bp
* 33432 34153: contig of 722 bp in length
* 34154 34253: gap of 100 bp
* 34254 34964: contig of 714 bp in length
* 34965 35064: gap of 100 bp
* 35065 35759: contig of 695 bp in length
* 35760 35859: gap of 100 bp
* 35860 36566: contig of 707 bp in length
* 36567 36666: gap of 100 bp
* 36667 37385: contig of 719 bp in length
* 37386 37485: gap of 100 bp
* 37486 38202: contig of 717 bp in length
* 38203 38302: gap of 100 bp
* 38303 39019: contig of 717 bp in length
* 39020 39119: gap of 100 bp
* 39120 39833: contig of 714 bp in length
* 39834 39933: gap of 100 bp
* 39934 40657: contig of 724 bp in length
* 40658 40757: gap of 100 bp
* 40758 41471: contig of 714 bp in length
* 41472 41571: gap of 100 bp
* 41572 42284: contig of 713 bp in length
* 42285 42384: gap of 100 bp
* 42385 43100: contig of 716 bp in length
* 43101 43200: gap of 100 bp
* 43201 43926: contig of 726 bp in length
* 43927 44026: gap of 100 bp
* 44027 44745: contig of 719 bp in length
* 44746 44845: gap of 100 bp
* 44846 45552: contig of 707 bp in length
* 45553 45652: gap of 100 bp
* 45653 46336: contig of 684 bp in length
* 46337 46436: gap of 100 bp
* 46437 47128: contig of 692 bp in length
* 47129 47228: gap of 100 bp
* 47229 47951: contig of 723 bp in length
* 47952 48051: gap of 100 bp
```

Alignment Scores:

```
Query No.: 4.46e-06 Length: 68314
Percent Similarity: 114.00 Matches: 22
Best Local Similarity: 83.87% Conservative: 4
Query Match: 70.97% Mismatches: 5
Indels: 0
Gaps: 0
```

US-09-513-999c-7869_copy_1_37 (1-37) x AC126345 (1-68314)

QY 6 AAlaLeuGlaAspSerPheSerLeuGlnGlyLeuLeuGlyProGluTyrValIysIeu 25

Db 43053 GCCCTGCGCTGAGTTGCACACAGATTGTGGGATCCCTTGGGCCGACGATTGTATAAGCTC 42994

QY 26 LeuGlyLeuCysValCysIeuSerGlyCysSer 36

Db 42993 CTGGGCTCTGTGTGTGTGTGACGACGCTCTCT 42961

RESULT 13

AC069245

LOCUS AC069245 143717 bp DNA linear HTG 09-MAY-2002

DEFINITION Homo sapiens chromosome 12 clone RP11-757N13, WORKING DRAFT

AC069245

AC069245.10 GI:20335692

KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.

SOURCE Homo sapiens.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 143717)
AUTHORS Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C., Alshrocks,S.L., Amaralunge,H.C., Are,J.R., Ayele,M., Banks,T., Barbarella,J., Benton,J., Bimaga,K., Blankenburg,K., Bonnin,D., Bouck,J., Bowles,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C., Burck,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R., Chen,Z., Chowdhry,I., Christopoulos,C., Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O., Denn,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H., Dugan-Rocha,S., Dublin,K.J., Barnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K., Harris,C., Harris,K., Hart,M., Havlak,P., Hawes,A., Hernandez,J., Hernandez,O., Hodgson,A., Hogues,M., Holloway,C., Hollins,B., Homs,F., Howard,S., Huber,J., Hulyk,S., Hume,J., Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudan,S., Karlsson,E., Kelly,S., Khan,U., King,L., Kovach,J., Kovar,C., Kratovic,J., Kureshi,A., Landry,N., Leal,B., Lewis,L.C., Lewis,L., Li,J., Li,Z., Licharge,O., Lieu,C., Liu,J., Liu,W., Louseged,H., Lozado,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J., Maheshwari,M., Mapua,P., Martin,R., Martindale,A., Martinez,E., Massey,E., Mawlinney,E., McLeod,M.P., Meador,M., Mel,G., Metzger,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K., Morgan,M., Morris,S., Moser,M., Neal,D., Newton,J., Mohabbat,K., Morgan,M., Morris,S., Nguyen,N., Nickerson,E., Nwokoko,S., Oguh,M., Okunolu,G., Oragunye,N., Oyiedo,R., Pace,A., Payton,B., Peery,J., Perez,L., Peters,L., Pickens,R., Primus,E., Pu,L.L., Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojudoan,I., Rolfe,M., Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shooshari,N., Sisson,I., Sodergren,E., Sonaike,T., Sparks,A., Stanley,H., Stone,H., Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H., Tansley,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S., Usmani,K., Vasquez,L., Vera,Y., Villalón,D., Vinson,R., Wang,Q., Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S., Williams,G., Williamson,A., Wleczky,R., Wooten,S., Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D., Weinstein,G., and Gibbs,R.

TITLE

JOURNAL

REFERENCE

AUTHORS

JOURNAL

REFERENCE

AUTHORS

JOURNAL

REFERENCE

AUTHORS

JOURNAL

REFERENCE

AUTHORS

JOURNAL

REFERENCE

AUTHORS

JOURNAL

Unpublished
2 (bases 1 to 143717)
Worley,K.C.
Direct Submission
Submitted (22-MAY-2000) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 143717)
Worley,K.C.
Direct Submission
Submitted (09-MAY-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Apr 28, 2002 this sequence version replaced gi:18449608.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HBX
Genetic clone name: RP11-757N13
----- Summary Statistics
Sequencing vector: M13
Chemistry: Dye-primer Bodipy: 19% of reads
Chemistry: Dye-terminator Big Dye: 81% of reads
Assembly program: Phrap; version 0.990329


```
* 3710 3809: gap of 100 bp
* 3810 6730: contig of 2921 bp in length
* 6731 6830: gap of 100 bp
* 6831 9206: contig of 2376 bp in length
* 9207 9306: gap of 100 bp
* 9307 12163: contig of 2857 bp in length
* 12164 12265: gap of 100 bp
* 12266 15381: contig of 3118 bp in length
* 15382 15481: gap of 100 bp
* 15482 20100: contig of 4619 bp in length
* 20101 20200: gap of 100 bp
* 20201 23885: contig of 3685 bp in length
* 23886 23985: gap of 100 bp
* 23986 27226: contig of 3241 bp in length
* 27227 27326: gap of 100 bp
* 27327 32869: contig of 5543 bp in length
* 32870 32969: gap of 100 bp
* 32970 36903: contig of 3934 bp in length
* 36904 37003: gap of 100 bp
* 37004 44368: contig of 7366 bp in length
* 44370 44469: gap of 100 bp
* 44470 53277: contig of 8808 bp in length
* 53278 53377: gap of 100 bp
* 53378 71089: contig of 17712 bp in length
* 71090 71189: gap of 100 bp
* 71190 90983: contig of 19794 bp in length
* 90984 91083: gap of 100 bp
* 91084 112306: contig of 21223 bp in length
* 112307 112406: gap of 100 bp
* 112407 152354: contig of 39948 bp in length.
Location/Qualifiers
1. 152354
```

```
FEATURES
Source
1. 152354
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-20p1"
/clone_id="RP11-11 Human Male BAC"
1. 1301
```

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

misc_feature

/note="assembly-fragment"

/note="assembly-fragment"

BASE COUNT 42989 a 30395 c 30652 g 46711 t 1607 others

ORIGIN

Alignment Scores:

Score: 9.89e-06

Percent Similarity: 114.00

Best Local Similarity: 85.71%

Query Match: 71.43%

DB: 59.07%

Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AC016472 (1-152354)

QY 9 AspSerPheSerSerLeuGlnGlyLeuGlnGlyProGlnTyValIysLeuGlnGlyLeu 28

DB 117786 GATGATTTCCACCTGCTGCTGATCTCGGACACAGATATTAACATCATGCTCTC 117845

QY 29 CysValCysLeuSerGlyCysSer 36

DB 117846 TGTATGTCCTGACGATGCTCT 117869

RESULT 15

AC027295

LOCUS

DEFINITION

Homo sapiens chromosome 12 clone RP11-293C20, WORKING DRAFT

SEQUENCE, 13 unordered pieces.

AC027295

VERSION

AC027295.15 GI:21431081

KEYWORDS

HTG: HTGS_PHASE1; HTGS_DRAFT.

SOURCE

human.

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.

I (bases 1 to 160066)

Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-osman,F.R., Allen,C.,

Alsbrooks,S.L., Amaralunge,H.C., Are,J.R., Ayalew,S.R., Banks,T.,

Barbieri,J., Benton,J., Bimaye,K., Blankenburg,K., Bonnin,D.,

Bouck,J., Bowler,S., Brileva,M., Brown,E., Brown,M., Bryant,N.P.,

Buhay,C., Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C.,

Carron,T.F., Carter,M., Cavazos,S.R., Chacko,J., Chavez,D.,

Chen,G., Chen,R., Chen,Z., Chowdhry,I., Christopoulos,C.,

Cleveland,C.D., Cox,C., Coyle,M.D., Dathorne,S.R., David,R.,

Davila,M.L., Davis,C., Davy-Carroll,L., Dederich,D.A.,

Delaney,K.R., Delgado,O., Dunn,A.L., Ding,Y., Dinh,H.H.,

Douthwaite,K.J., Draper,H., Dugan-Hochs,S., Durkin,J.,

Earnhart,C., Edgar,D., Edwards,C.C., Elhaj,C., Escotto,M.,

Falls,T., Ferraguto,D., Flagg,N., Ford,J., Foster,P., Frantz,P.,

Gabisi,A., Gao,J., Garcia,A., Garner,T., Garza,N., Gill,R.,

Garratt,J., Guevara,W., Gunaratne,P., Hale,S., Hamilton,K.,

Harris,C., Harris,K., Hart,M., Havlak,P., Hawes,A., Hernandez,J.,

Hernandez,O., Hodgson,A., Hogues,M., Holloway,C., Hollins,B.,

Honsi,F., Howard,S., Huber,J., Huijk,S., Hume,J., Jackson,L.E.,

Jacobson,B., Jia,Y., Johnson,R., Jolivet,S., Joudah,S.,

Karlsson,E., Kelly,S., Khan,U., King,L., Korah,J., Kovar,C.,

Kratovic,J., Kureshi,A., Landry,N., Deal,B., Lewis,L.C., Lewis,L.,

Li,J., Li,Z., Licharge,O., Lieu,C., Liu,J., Liu,W., Lounsbury,H.,

Lozano,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R., Ma,J.,

Maneshwari,M., Mapua,P., Martin,R., Martindale,A., Martinez,E.,

Massey,E., Mawhney,E., McLeod,M.P., Meador,M., Mei,G., Metzger,M.,

Miner,G., Miner,Z., Mitchell,T., Mohabat,K., Morgan,M., Morris,S.,

Moser,M., Neal,D., Newton,J., Newton,N., Nguyen,A., Nguyen,N.,

Nguyen,N., Nickerson,E., Nwokenwo,S., Ogih,M., Okwouu,G.,

Oragunye,N., Oviedo,R., Pace,A., Payton,B., Peery,D., Perez,L.,

Peters,L., Pickens,R., Primus,E., Pu,L.L., Quiles,M., Ren,Y.,

Rivera,M., Rojas,A., Rojuben,I., Rolle,M., Ruiz,S., Saverly,G.,

Scherer,S., Scott,G., Shen,H., Shoshchari,N., Sisson,I.,

Sodergren,E., Sonalke,T., Sparks,A., Stanley,R., Stone,H.,

Sutton,A., Sytek,A., Tabor,P., Tamerisa,A., Tamerisa,K., Tang,H.,

Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N., Thomas,S.,

Usmanli,K., Vasquez,B., Vera,V., Villalobon,D., Vinson,K., Wang,Q.,

Wang,S., Ward-Moore,S., Warren,R., Washington,C., Watlington,S.,

Williams,G., Williamson,A., Wleczky,R., Wooden,S., Worley,K.,

Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Nelson, D.,
Weinstock, G. and Gibbs, R.
TITLE Direct Submission
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 160066)
AUTHORS Worley, K.C.
TITLE Direct Submission
JOURNAL Submitted (30-MAR-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
3 (bases 1 to 160066)
Worley, K.C.
TITLE Direct Submission
JOURNAL Submitted (18-JUN-2002) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Jun 17, 2002 this sequence version replaced gi:20335550.

COMMENT

----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

Web site: <http://www.hgsc.bcm.tmc.edu/>

Drafting Center Code: BCM

Contact: hgsc-help@bcm.tmc.edu

----- Project Information

Center project name: HARS

Center clone name: RP11-293C20

----- Summary Statistics

Sequencing vector: M13;

Chemistry: Dye-primer Bodipy: 42% of reads

Chemistry: Dye-terminator Big Dye: 58% of reads

Assembly program: Phrap; version 0.990329

Consensus quality: 144209 bases at least Q40

Consensus quality: 149815 bases at least Q30

Consensus quality: 153559 bases at least Q20

Estimated insert size: 1574/3; sum-of-coverage estimation

Quality coverage: 4x in Q20 bases; sum-of-coverage estimation

* NOTE: Estimated insert size may differ from sequence length

* (see http://www.hgsc.bcm.tmc.edu/docs/genbank_draft_data.html).

* NOTE: This is a 'working draft' sequence. It currently

* consists of 13 contigs. The true order of the pieces

* is not known and their order in this sequence record is

* arbitrary. Gaps between the contigs are represented as

* runs of N, but the exact sizes of the gaps are unknown.

* This record will be updated with the finished sequence

* as soon as it is available and the accession number will

* be preserved.

* 1 3659: contig of 3659 bp in length

* 3660 3759: gap of unknown length

* 3760 5914: contig of 2155 bp in length

* 5915 6014: gap of unknown length

* 6015 9826: contig of 3812 bp in length

* 9827 9926: gap of unknown length

* 9927 12721: contig of 2794 bp in length

* 12721 12820: gap of unknown length

* 12820 20586: contig of 7766 bp in length

* 20587 20687: gap of unknown length

* 20687 27322: contig of 6636 bp in length

* 27323 27422: gap of unknown length

* 27423 39529: contig of 12107 bp in length

* 39530 39629: gap of unknown length

* 39630 54210: contig of 14581 bp in length

* 54211 54310: gap of unknown length

* 54311 64148: contig of 9838 bp in length

* 64149 64248: gap of unknown length

* 64249 79424: contig of 15176 bp in length

* 79425 79524: gap of unknown length

* 79525 99543: contig of 20019 bp in length

* 99544 99644: gap of unknown length

* 99645 125583: contig of 25939 bp in length

* 125583 125683: gap of unknown length

* 125683 160066: contig of 34384 bp in length

FEATURES

Location/Qualifiers

source

1..160066

/organism="Homo sapiens"

/db_xref="taxon:9606"

/chromosome="12"

/clone="RP11-293C20"

BASE COUNT 47149 a 32911 c 32401 g 46353 t 1252 others

ORIGIN

Alignment Scores:

Pred. No.:

Score:

Percent Similarity:

Best Local Similarity:

Query Match:

DB:

1.04e-05

114.00

85.71%

71.43%

59.07%

2

Length:

Matches:

Conservative:

Mismatches:

Indels:

Gaps:

160066

20

4

4

0

0

US-09-513-999c-7869_COPY_1_37 (1-37) x AC027295 (1-160066)

QY 9 ApsSerPheSerSerLeuGlnGlyLeuLeuGlyProGluTrpValLysLeuLeuGlyLeu 28

DB 49872 GATGGATTTCACCTCTGCGATCCTCGGACACAGTATGTAACATCATGCGTCTC 49931

QY 29 CysValCysLeuSerGlyCysSer 36

DB 49932 TGTATGTGCTGTCAGCAGATGCTCT 49955

Search completed: April 24, 2003, 23:01:33

Job time : 1526 secs

548.184 Million cell updates/sec

Sequence: 1 MGGSFALQDSFSSLQGLGPEYVKLLGLCVCLSGCST 37

Xgapop	10.0	Xgapext	0.5
Ygapop	10.0	Ygapext	0.5
Fgapop	6.0	Fgapext	7.0
Delop	6.0	Delext	7.0

Total number of hits satisfying chosen parameters: 4370478

Post-processing: Minimum match 0%

Listing first 45 summaries

Command line parameters:

```
MODEL=irmme_p2n.model -DEV=xlh
-Q/cgml2/USPTO.spool/US0951999/r/unat.18042003.170936.28347/app.quick.fasta_1.199
-DB=n.GeneSeq.101002 -QEM=fastap -SUFFIX=p2n.rng -MINMATCH=0.1 -LOOCL=0
-LOOEXT=0 -UNITS=bits -START=1 -END=-1 -MATRIX=blotsum62 -TRANS=human400.cdi
-LIST=45 -DOCLATION=200 -THR_SCORE=ptc -THR_MAX=100 -THR_MIN=0 -ALGIN=15
-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZ=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09513999.ecgn1.p1.2006.atunat.18042003.170936.28347 -NCPU=6 -ICPU=3
-NO_XLPHY -NO_MAMP -LARGEQUERY -NEG_SCORES=0 -WAIT -LONCLONG -DEV_TIMECOUT=120
-WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6 -FGAPEXT=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXEXT=7
```

Database :

1:	/SID2/gcgdata/genseq/genseqn-emb1/NA1980.DAT *
2:	/SID2/gcgdata/genseq/genseqn-emb1/NA1981.DAT *
3:	/SID2/gcgdata/genseq/genseqn-emb1/NA1982.DAT *
4:	/SID2/gcgdata/genseq/genseqn-emb1/NA1983.DAT *
5:	/SID2/gcgdata/genseq/genseqn-emb1/NA1984.DAT *
6:	/SID2/gcgdata/genseq/genseqn-emb1/NA1985.DAT *
7:	/SID2/gcgdata/genseq/genseqn-emb1/NA1986.DAT *
8:	/SID2/gcgdata/genseq/genseqn-emb1/NA1987.DAT *
9:	/SID2/gcgdata/genseq/genseqn-emb1/NA1988.DAT *
10:	/SID2/gcgdata/genseq/genseqn-emb1/NA1989.DAT *
11:	/SID2/gcgdata/genseq/genseqn-emb1/NA1990.DAT *
12:	/SID2/gcgdata/genseq/genseqn-emb1/NA1991.DAT *
13:	/SID2/gcgdata/genseq/genseqn-emb1/NA1992.DAT *
14:	/SID2/gcgdata/genseq/genseqn-emb1/NA1993.DAT *
15:	/SID2/gcgdata/genseq/genseqn-emb1/NA1994.DAT *
16:	/SID2/gcgdata/genseq/genseqn-emb1/NA1995.DAT *
17:	/SID2/gcgdata/genseq/genseqn-emb1/NA1996.DAT *
18:	/SID2/gcgdata/genseq/genseqn-emb1/NA1997.DAT *
19:	/SID2/gcgdata/genseq/genseqn-emb1/NA1998.DAT *
20:	/SID2/gcgdata/genseq/genseqn-emb1/NA1999.DAT *
21:	/SID2/gcgdata/genseq/genseqn-emb1/NA2000.DAT *
22:	/SID2/gcgdata/genseq/genseqn-emb1/NA2001A.DAT *
23:	/SID2/gcgdata/genseq/genseqn-emb1/NA2001B.DAT *
24:	/SID2/gcgdata/genseq/genseqn-emb1/NA2002.DAT *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed and is derived by analysis of the total score distribution.

	Result No.	Score	Query Match	Length	DB	ID	Description
ALIGNMENTS	1	193	100.0	447	21	AAC03794	Human secreted protein
	2	193	100.0	447	21	AAZ24680	Human 5' EST isolate
	3	95	45.9	128600	22	ABK83461	Human cDNA difference
	4	88.5	45.9	570	22	ABA63453	Human foetal liver
	5	88.5	45.9	570	22	ABA30652	Probe #918 for gene
	6	88.5	45.9	570	22	AAI11985	Human brain expression
	7	88.5	45.9	570	22	AAK37688	Human bone marrow
	8	88.5	45.9	570	22	AAI16447	Human bone marrow
	9	88.5	45.9	570	22	AAI14563	Probe #8380 for gene
	10	88.5	45.9	570	22	AAI14563	Probe #12249 used
	11	88	45.6	1982	21	ABS11680	Human genome-derived
	12	86	44.6	1299	23	AAK68089	Human secreted protein
	13	85	44.0	1299	23	ASNS91827	DNA encoding novel
	14	83	43.0	1909	23	AAK88434	Human ORFX polynucleotide
	15	83	43.0	1909	23	AAK88434	DNA encoding novel
	16	78	40.4	609	22	AAH96980	DNA encoding novel
	17	71	36.8	11534	24	AAH65888	Novel Human polynucleotide
	18	69	35.8	2787	23	AAK68019	Human immune system
	19	69	35.8	2787	23	AAK68019	DNA encoding novel
	20	67	34.7	1996	22	AAI16896	DNA encoding novel
	21	65	33.7	447	22	AAI16101	Human nervous system
	22	65	33.7	447	22	AAI24945	Human breast cancer
	23	64	33.2	36901	20	AAZ23892	Human breast cancer
	24	64	33.2	38866	20	AAZ23892	Murine LOBO genome
	25	63	32.6	162	22	AAW57562	Murine LOBO homologue
	26	63	32.6	519	22	ABA63320	Human foetal liver
	27	60.5	31.3	2730	22	AAK84293	Human EXCS encoding
	28	60.5	31.3	2453	22	AAK27689	DNA encoding novel
	29	59	30.6	481	23	AAV05461	Human prostate expression
	30	59	30.6	560	22	AAK78945	Human prostate expression
	31	58.5	30.3	27666	23	ABL08332	Human immune/haematoma
	32	58	30.1	2085	20	AAAX01362	Drosophila melanogaster
	33	57.5	29.8	1605	24	ABL40748	Nucleobase permease
	34	57	29.5	273	22	AAK86888	Chicken heparanase
	35	57	29.5	7379	19	AAV49653	Human immune/haematoma
	36	56.5	29.3	259	23	AAV05218	Human SCL DNA. Homo sapiens
	37	56.5	29.3	229	23	ABY1487	Human prostate expression
	38	56.5	29.3	400	23	ABY3471	Human prostate expression
	39	56.5	29.3	650	24	ABO56543	Human colon cancer
	40	56.5	29.3	2076	22	AAK54519	DNA encoding novel
	41	56	29.0	1226	22	AAK53271	Human polynucleotide
	42	56	29.0	1380	24	AAK7394	Bacillus licheniformis
	43	56	29.0	2015	22	AAH57417	Human stomach cell
	44	56	29.0	2073	20	AAK87925	Human protease HPP
	45	56	29.0	2118	22	AAI60469	Human polynucleotide
ALIGNMENTS							
RESULT 1							
AAAC03794							
ID	AAAC03794	standard;	cDNA;	447	BP.		
XX	AAAC03794;						
XX	AC						
DT	06-Oct-2000	(first entry)					
XX	Human	secreted protein	5'	EST,	SEQ ID NO:	3792.	
DE	Human	secreted protein	5'	EST,	SEQ ID NO:	3792.	
XX	Human	5' EST; expressed sequence tag; secreted protein; cDNA isolation;					
KW	gene therapy; chromosome mapping; ss.						
KW	Homo sapiens.						
OS	Homo sapiens.						
XX	EPI033401-A2.						
FN	EPI033401-A2.						
XX	06-SEP-2000.						

KW cardiac reperfusion injury; renal reperfusion injury; ARDS;
 KW adult respiratory distress syndrome; inflammatory bowel disease;
 KW Crohn's disease; ulcerative colitis; periodontal disease;
 KW granulocyte activation; chronic inflammation; allergy.
 OS
 OS Homo sapiens.
 PN W0200228999-A2.
 PD 11-APR-2002.
 XX 03-OCT-2001; 2001MO-US30821.
 PF 03-OCT-2001; 2000US-237189P.
 PR 03-OCT-2000; 2000US-237189P.
 XX
 XX (GENE-) GENE LOGIC INC.
 PA
 XX
 XX Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;
 P1
 XX WPI: 2002-435328/46.
 DR
 PT Detecting granulocyte activation by detecting differential expression
 PT of genes associated with granulocyte activation, which serves as
 PT diagnostic markers that is useful for monitoring disease states and
 PT drug toxicity -
 XX
 XX
 PS Claim 1; SEQ ID NO 32; 114pp; English.
 CC
 CC The invention relates to detecting (M1) granulocyte (GC) activation
 CC (GCA), by detecting the level of expression of gene(s) (Gs) identified by
 CC DNA chip analysis as given in the specification, and comparing
 CC the expression level to an expression level in an unactivated
 CC GC, where differential expression of Gs is indicative of GCA.
 CC Also included are modulating (M2) GA by contacting GC with an agent
 CC that alters the expression of at least one gene in Gs; (2) screening (M3)
 CC for an agent capable of modulating GCA or an inflammation (especially
 CC chronic) in a tissue, an allergic response in a subject, exposure of a
 CC subject to a pathogen or sterile inflammatory disease using the
 CC gene expression profile; (3) detecting (M4) an inflammation (especially
 CC chronic) in a tissue, an allergic response in a subject, exposure of a
 CC subject to a pathogen or sterile inflammatory disease, by detecting the
 CC level of expression in a sample of the tissue of gene(s) from Gs, where
 CC the level of expression of the gene is indicative of inflammation;
 CC (4) treating (M5) an inflammation (especially chronic) or in a tissue,
 CC an allergic response in a subject, exposure of a subject to a pathogen
 CC or sterile inflammatory disease, by contacting a tissue having
 CC inflammation with an agent that modulates the expression of gene(s)
 CC from Gs in the tissue. M1 is useful for detecting GCA. M2 is useful for
 CC modulating GA; M3 is useful for screening an agent capable of modulating
 CC GCA preferably in an inflammation in a tissue; M4 is useful for
 CC detecting an inflammation (especially chronic) in a tissue, an allergic
 CC response in a subject, exposure of a subject to a pathogen or sterile
 CC inflammatory disease (e.g. psoriasis, rheumatoid arthritis,
 CC glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal
 CC reperfusion injury, ARDS, adult respiratory distress syndrome,
 CC inflammatory bowel disease, Crohn's disease, ulcerative colitis,
 CC periodontal disease; also bacterial infection, viral infection,
 CC parasitic infection, protozoal infection, fungal infection and M5 is
 CC useful for treating one of the above conditions. The present
 CC sequence represents a gene differentially expressed in granulocytes.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX
 SO Sequence 128600 BP; 36139 A; 24970 C; 26316 G; 41175 T; 0 other;

Alignment Scores:
 Pred. NO.: 0.0459 Length: 128600
 Score: 95.00 Matches: 21
 Percent Similarity: 64.864 Conservative: 3
 Best Local Similarity: 56.768 Mismatches: 9
 Query Match: 49.224 Indels: 4

```

DB:                               24                               Gaps:                               1
US-09-513-999C-7869_COPY_1_37 (1-37) x ABA63451 (1-128600)
Oy      3  GlycerPheAlaLeuGlnAspSerPheSerSerLeuGln-----GlyLeuLeu 18
          |||:::                               ::|  |||
Db 126328 GGGGCTATGCTGTGACCTCCACCTTGACTTCATTGCAAGACGATTTGTGGGGCTCTG 126269
Oy      19  GlProGluTyrValLysLeuLeuGlyLeuCySValLysLeuSerGlyCyS 35
          |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db 126268 GGGCTGAGGTATGTAAAGCTCTCGGCTCTGTGCATGCCTGACGAGCTGC 126218

RESULT 4
ABA63453/c
ID      ABA63453 standard; DNA: 570 BP.
XX
XX      ABA63453;
XX
DT      01-FEB-2002 (first entry)
XX
XX      Human foetal liver single exon nucleic acid probe #11758.
DE
XX      Human; foetal liver; gene expression; single exon nucleic acid probe; ss
XX
XX      Homo sapiens.
OS
XX      WO200157277-A2.
PN
PD      09-AUG-2001.
XX
XX      30-JAN-2001; 2001WO-0500669.
PF
PR      04-FEB-2000; 2000US-0180312.
PR      26-MAY-2000; 2000US-0207456.
PR      30-JUN-2000; 2000US-0608408.
PR      03-AUG-2000; 2000US-0632386.
PR      21-SEP-2000; 2000US-0234687.
PR      27-SEP-2000; 2000US-0236359.
PR      04-OCT-2000; 2000GB-0024263.
XX
XX      (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX      Penn SG, Hanzel DK, Chen W, Rank DR;
PI
DR      WPI: 2001-483447/52.
XX
XX      Human genome-derived single exon nucleic acid probes useful for
PT      analyzing gene expression in human fetal liver -
PS      Claim 1; SEQ ID NO 11758; 6399p + sequence listing; English.
XX
XX      The invention relates to a single exon nucleic acid probe for
CC      measuring human gene expression in a sample derived from human foetal
CC      liver. The single exon nucleic acid probes may be used for predicting,
CC      measuring and displaying gene expression in samples derived from human
CC      fetal liver. The present sequence is a single exon nucleic acid
CC      probe of the invention.
CC      Note: The sequence data for this patent did not form part of the
CC      printed specification, but was obtained in electronic format directly
CC      from WIPO at ftp.wipo.int/pub/published_pat_sequences.
XX
XX      Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Alignment Scores:
Pred. No.:      0.000408      Length:      570
Score:          88.50         Matches:      21
Percent Similarity: 61.11%     Conservative: 1
Best Local Similarity: 58.33%   Mismatches: 7
Query Match:    45.85%         Indels:      7
DB:            22             Gaps:        1
US-09-513-999C-7869_COPY_1_37 (1-37) x ABA63453 (1-570)

```

```

Oy      1 MetGtGcSerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyPro 20
Db      568 ATGAATGATCT-----CTGCTTGTGCGGATTCCTTGCGCTG 530
Oy      21 GluTyValLysLeuLeuGlyLeuCysValCysLeuSerGlyCysSer 36
Db      529 GAGTATGTAATAATTCCTGGCTTGTGTGTGTGCTGAGTGCGCGCTCT 482

RESULT 5
ABA30652/c
ID      ABA30652 standard; DNA: 570 BP.
XX
XX      ABA30652;
AC
XX
XX      23-JAN-2002 (first entry)
DE
XX      Probe #9118 for gene expression analysis in human heart cell sample.
XX      Human; gene expression; heart; microarray; vascular system; probe;
XX      cardiovascular disease; hypertension; cardiac arrhythmia;
XX      congenital heart disease; ss.
XX      Homo sapiens.
XX
XX      WO200157274-A2.
XX
XX      09-AUG-2001.
XX
XX      30-JAN-2001: 2001WO-US00666.
XX
XX      04-FEB-2000: 2000US-0180312.
XX      26-MAY-2000: 2000US-0207456.
XX      30-JUN-2000: 2000US-0608408.
XX      03-AUG-2000: 2000US-0632366.
XX      21-SEP-2000: 2000US-0234687.
XX      27-SEP-2000: 2000US-0236359.
XX      04-OCT-2000: 2000GB-0024263.
XX
XX      (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX      Penn SG, Hanzel DK, Chen W, Rank DR;
XX
XX      WPI; 2001-48889/53.
XX
XX      Single exon nucleic acid probes for analyzing gene expression in human
XX      hearts -
XX
XX      Claim 1; SEQ ID No 9118; 530bp; English.

The present invention relates to single exon nucleic acid probes for
measuring human gene expression in a sample derived from human heart. The
present sequence is one such probe. The probes may be used for
predicting, measuring and displaying gene expression in samples derived
from the human heart via microarrays. By measuring gene expression, the
probes are useful for predicting, diagnosing, grading, staging,
monitoring and prognosing diseases of the human heart and vascular system
e.g. cardiovascular disease, hypertension, cardiac arrhythmias and
congenital heart disease.
Note: The sequence data for this patent did not form part of the printed
specification, but was obtained in electronic format directly from WIPO
at ftp.wipo.int/pub/published_pct-sequences.
XX
XX      Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
XX

Alignment Scores:
Pred. No.:      0.00408      Length:      570
Score:          88.50       Matches:      21
Percent Similarity: 61.11%   Conservative: 1
Best local Similarity: 58.33%   Mismatches: 7
Query Match:    45.85%      Indels:      7
DB:             22          Gaps:        1

(US-09-513-999C-7869-COPY_1-37 (1-37) x ABA30652 (1-570))

```

Oy		1 MetGtGlySerPheAlaLeuGlInAspSerPheSerIeuGlnGlyLeuLeugLyPro	20
Dd	568 ATGAATGCATCT-----CCTGCCITTCGGATTCCTTGGCGTc	530	
Oy		21 GlutryValLysLeuLeuGlyLeuCysValCyslserGlyCysser	36
Dd	529 GAGTATGTAAATAATTCCTGGTGCTTGTGTGTGCCTGAGTAGCGCGCTCT	482	

RESULT 6

AAK11985/c
 ID AAK11985 standard; DNA; 570 BP.
 xx
 AC AAK11985;
 xx
 DT 05-NOV-2001 (first entry)
 DE Human brain expressed single exon probe SEQ ID NO: 11976.
 XX
 KW Human: brain expressed exon; gene expression analysis; probe;
 KM microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
 KK epilepsy; cancer; SS.
 OS Homo sapiens.
 XX
 PN WO200157275-A2.
 XX
 PD 09-AUG-2001.
 PF 30-JAN-2001; 2001WO-USO0667.
 PR 04-FEB-2000; 2000US-0180312.
 PR 26-MAY-2000; 2000US-0207456.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0632366.
 PR 21-SEP-2000; 2000US-0234687.
 PR 27-SEP-2000; 2000US-0236359.
 PR 04-OCT-2000; 2000GB-0024263.
 XX
 PA (MOLE-) MOLECULAR DYNAMICS INC.
 PI Penn SG, Hanzel DK, Chen W, Rank DR:
 XX
 DR WPJ; 2001-483446/52.

Single exon nucleic acid probes for analyzing gene expression in human brains -

Example 4; SEQ ID NO: 11976; 650bp + Sequence listing; English.

The present invention provides a number of single exon nucleic acid probes which are derived from genomic sequences expressed in the human brain. They can be used to measure gene expression in brain cell samples which may enable the diagnosis and improved treatment of nervous system diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia, CC epilepsy and cancers. The present sequence is one of the probes of the invention.

SEQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other:

Alignment Scores:			
Pred. NO.: 0.000408	Length: 570		
Score: 88.50	Matches: 21		
Percent Similarity: 61.1%	Conservative: 1		
Best Local Similarity: 58.33%	Mismatches: 7		
Query Match: 45.85%	Indels: 7		
DB: 22	Gaps: 1		

```

US-09-513-999C-7869_COPY_1_37 (1-37) x AAK11985 (1-570)
OY      1 MetGtGlySerPheAlaLeuGlInAspSerPheSerIeuGlnGlyLeuLeugLyPro 20
       |||         |||||          :|||   |||   ||||||
Dd     568 ATGAATGCATCT-----CCTGCCITTCGGATTCCTTGGCGTc 530

```

OY 21 GUTYrVallylsleuLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36
 ||||||| ||||||| ||||||| ||||||| |||
 Db 529 GAGTATGTAAATTCCTGGGCTTTGTGTGTGCTGAGGCGGCTCT 482

RESULT 7

AAK37688/c
 ID AAK37688 standard; DNA; 570 BP.

AC AAK37688;

DT 06-NOV-2001 (first entry)

DE Human bone marrow expressed single exon probe SEQ ID NO: 12245.

KW Human; bone marrow expressed exon; gene expression analysis; probe;
 microarray; cancer; leukemia; lymphoma; myeloma; ss.

OS Homo sapiens.

WO200157276-A2.

PD 09-AUG-2001.

PE 30-JAN-2001; 2001WO-US00668.

PR 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

PI Penn SG, Hanzel DK, Chen W, Rank DR;

DR WPI: 2001-488900/53.

XX Human genome-derived single exon nucleic acid probes useful for

PT analyzing gene expression in human bone marrow -

XX Example 4; SEQ ID NO: 12245; 658bp + Sequence Listing; English.

CC The present invention provides a number of single exon nucleic acid

CC probes which are derived from genomic sequences expressed in the human

CC bone marrow. They can be used to measure gene expression in bone marrow

CC samples, which may enable the improved diagnosis and treatment of cancers

CC such as lymphoma, leukemia and myeloma. The present sequence is one of

XX the probes of the invention.

SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Alignment Scores:

Pred. No.: 0.000408 Length: 570

Score: 88.50 Matches: 21

Percent Similarity: 61.11% Conservative: 1

Best Local Similarity: 58.33% Mismatches: 7

Query Match: 45.85% Indels: 7

DB: Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x AAK37688 (1-570)

OY 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuLeuGlyPro 20

Db 568 ATGAATGATCT-----CCTGCTTCTGGGATTCTTGGGCTG 530

OY 21 GUTYrVallylsleuLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36

Db 529 GAGTATGTAAATTCCTGGGCTTTGTGTGTGCTGAGGCGGCTCT 482

RESULT 8

AA118447/c
 ID AA118447 standard; DNA; 570 BP.

AC AA118447;

DT 12-OCT-2001 (first entry)

DE Probe #8380 for gene expression analysis in human cervical cell sample.

KW Probe; human; microarray; gene expression; cervical epithelial cell;
 cervical cancer; ss.

OS Homo sapiens.

WO200157278-A2.

PD 09-AUG-2001.

PE 30-JAN-2001; 2001WO-US00670.

PR 04-FEB-2000; 2000US-0180312.

PR 26-MAY-2000; 2000US-0207456.

PR 30-JUN-2000; 2000US-0608408.

PR 03-AUG-2000; 2000US-0632366.

PR 21-SEP-2000; 2000US-0234687.

PR 27-SEP-2000; 2000US-0236359.

PR 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

PI Penn SG, Hanzel DK, Chen W, Rank DR;

DR WPI: 2001-488901/53.

XX Human genome-derived single exon nucleic acid probes useful for

PT analyzing gene expression in human cervical epithelial cells -

XX Claim 25; SEQ ID No 8380; 487bp; English.

CC The present invention relates to human single exon nucleic acid probes

CC (SNP). The present sequence is one such probe. The SNPs are derived

CC from human HeLa cells. The SNPs can be used to produce a single exon

CC microarray, which can be used for measuring human gene expression in a

CC sample derived from human cervical epithelial cells. By measuring gene

CC expression, the probes are therefore useful in grading and/or staging

CC of diseases of the cervix, notably cervical cancer.

CC Note: The sequence data for this patent did not form part of the printed

CC specification, but was obtained in electronic format directly from WIPO

XX at ftp.wipo.int/pub/published_pct_sequences.

SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Alignment Scores:

Pred. No.: 0.000408 Length: 570

Score: 88.50 Matches: 21

Percent Similarity: 61.11% Conservative: 1

Best Local Similarity: 58.33% Mismatches: 7

Query Match: 45.85% Indels: 7

DB: Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x AA118447 (1-570)

OY 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuLeuGlyPro 20

Db 568 ATGAATGATCT-----CCTGCTTCTGGGATTCTTGGGCTG 530

OY 21 GUTYrVallylsleuLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36

Db 529 GAGTATGTAAATTCCTGGGCTTTGTGTGTGCTGAGGCGGCTCT 482

RESULT 9

AA143563/c
 ID AA143563 standard; DNA; 570 BP.

```
XX AC AAI43563;
XX XX
XX DT 17-OCT-2001 (first entry)
XX DE Probe #12249 used to measure gene expression in human placenta sample.
XX XX
XX KW Probe; microarray: human; placenta; antenatal diagnosis;
XX KW genetic disorder; ss.
XX OS Homo sapiens.
XX PN WO200157272-A2.
XX PD 09-AUG-2001.
XX PF 30-JAN-2001; 2001WO-US00663.
XX PR 04-FEB-2000; 2000US-0180312.
XX PR 26-MAY-2000; 2000US-0207456.
XX PR 30-JUN-2000; 2000US-0608408.
XX PR 03-AUG-2000; 2000US-0632366.
XX PR 21-SEP-2000; 2000US-0234687.
XX PR 27-SEP-2000; 2000US-0236359.
XX PR 04-OCT-2000; 2000GB-0024263.
XX PA (MOLE-) MOLECULAR DYNAMICS INC.
XX PT Penn SG, Hanzel DK, Chen W, Rank DR;
XX DR WPI: 2001-488897/53.
XX XX
XX PT Human genome-derived single exon nucleic acid probes useful for
XX PT analyzing gene expression in human placenta -
XX PS Claim 25: SEQ ID No 12249; 654bp; English.
XX XX
XX CC The present invention relates to single exon nucleic acid probes (SENP).
XX CC The present sequence is one such probe. The probes are useful for
XX CC producing a microarray for predicting, measuring and displaying gene
XX CC expression in samples derived from human placenta. The probes are useful
XX CC for antenatal diagnosis of human genetic disorders.
XX SQ Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
XX
XX Alignment Scores:
XX Pred. No.: 0.000408 Length: 570
XX Pre: 88.50 Matches: 21
XX Percent Similarity: 61.11% Conservative: 1
XX Best Local Similarity: 58.33% Mismatch: 7
XX Query Match: 45.85% Indels: 7
XX DB: Gaps: 1
XX
XX US-09-513-999c-7869_COPY_1_37 (1-37) x AAI43563 (1-570)
XX
XX OY 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuLeuGlyPro 20
XX ||| |||||
XX ||| |||||
XX Db 568 ATGATGATCT-----CCTGCTTGCTGGGATTCCTTGGGCTG 530
XX
XX OY 21 GluTyrValIleuLeuGlyLeuGlyCysValGlyLeuGlyCysSer 36
XX ||||| ||||| ||||| ||||| ||||| |||||
XX ||||| ||||| ||||| ||||| ||||| |||||
XX Db 529 GAGTATGTAAATTCCTGGGCTTGTGTGCTGAGTGCCTCT 482
XX
XX RESULT 10
XX ABS11680/c
XX ID ABS11680 standard; DNA; 570 BP.
XX AC ABS11680;
XX XX
XX DT 19-AUG-2002 (first entry)
XX DE Human genome-derived single exon probe from lung SEQ ID No 11671.
XX XX
```

```
KW Human; ds: single exon probe; asthma; lung cancer; COPD; ILD;
KW chronic obstructive pulmonary disease; interstitial lung disease;
KW familial idiopathic pulmonary fibrosis; neurofibromatosis;
KW tuberous sclerosis; Gaucher's disease; Niemann-Pick disease;
KW Hermansky-Pudlak syndrome; sarcoidosis; pulmonary haemosiderosis;
KW pulmonary histiocytosis; lymphangioleiomyomatosis; Kargener syndrome;
KW pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
KW primary ciliary dyskinesia; pulmonary hypertension;
KW hyaline membrane disease.
XX
XX OS Homo sapiens.
XX PN WO200186003-A2.
XX PD 15-NOV-2001.
XX PF 30-JAN-2001; 2001WO-US00665.
XX PR 04-FEB-2000; 2000US-180312P.
XX PR 26-MAY-2000; 2000US-207456P.
XX PR 30-JUN-2000; 2000US-0608408.
XX PR 03-AUG-2000; 2000US-0632366.
XX PR 21-SEP-2000; 2000US-234687P.
XX PR 27-SEP-2000; 2000US-236359P.
XX PR 04-OCT-2000; 2000GB-0024263.
XX PA (MOLE-) MOLECULAR DYNAMICS INC.
XX PT Penn SG, Hanzel DK, Chen W, Rank DR;
XX DR WPI: 2002-114183/15.
XX XX
XX PT Spatially-addressable set of single exon nucleic acid probes, used to
XX PT measure gene expression in human lung samples -
XX PS Claim 1: SEQ ID No 11671; 634bp; English.
XX XX
XX CC The invention relates to a spatially-addressable set of single exon
XX CC nucleic acid probes for measuring gene expression in a sample derived
XX CC from human lung comprising single exon nucleic acid probes having one of
XX CC 12614 nucleic acid sequences mentioned in the specification, or their
XX CC complements or the 12387 open reading frames derived from the 12614
XX CC probes. Also included are a microarray comprising the novel set of
XX CC probes; the novel set of probes which hybridise at high stringency to a
XX CC nucleic acid expressed in the human lung; measuring gene expression in a
XX CC sample derived from human lung, comprising (a) contacting the array with
XX CC a collection of detectably labeled nucleic acids derived from human lung
XX CC mRNA, and (b) measuring the label detectably bound to each probe of
XX CC the array; identifying exons in a eukaryotic genome, comprising
XX CC (a) algorithmically predicting at least one exon from genomic sequences
XX CC of the eukaryote; and (b) detecting specific hybridisation of detectably
XX CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
XX CC having a fragment identical to the predicted exon, the probe is included
XX CC in the above mentioned microarray; assigning exons to a single gene,
XX CC comprising (a) identifying exons from genomic sequence by the method
XX CC above and (b) measuring the expression of each of the exons in several
XX CC tissues and/or cell types using hybridisation to a single exon
XX CC microarrays having a probe with the exon, where a common pattern of
XX CC expression of the exons in the tissues and/or cell types indicates that
XX CC the exons should be assigned to a single gene; a peptide comprising one
XX CC of 12011 sequences, mentioned in the specification, or encoded by the
XX CC probes/open reading frames (ORF). The probes are used for gene
XX CC expression analysis, and for identifying exons in a gene, particularly
XX CC using human lung derived mRNA and for the study of lung diseases
XX CC such as asthma, lung cancer, chronic obstructive pulmonary disease
XX CC (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
XX CC fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
XX CC Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
XX CC haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis,
XX CC pulmonary alveolar proteinosis, Kargener syndrome, fibrocystic
XX CC pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
XX CC and hyaline membrane disease. The present sequence is a single exon
XX CC probe of the invention.
```



```
PR 31-MAR-2000; 2000US-0540217.
PR 23-AUG-2000; 2000US-0649167.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Drmanac RT, Liu C, Tang YT;
XX
XX WPI: 2001-639362/73.
DR P-PSDB: AAB24247.
XX
XX New isolated polynucleotide and encoded polypeptides, useful in
PT diagnostics, forensics, gene mapping, identification of mutations
PT responsible for genetic disorders or other traits and to assess
PT biodiversity
XX
XX Claim 1: SEQ ID No 24238; 103pp; English.
XX
XX The invention relates to isolated polynucleotide (I) and
CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
CC and gene mapping, and in recombinant production of (II). The
CC polynucleotides are also used in diagnostics as expressed sequence tags
CC for identifying expressed genes. (I) is useful in gene therapy techniques
CC to restore normal activity of (II) or to treat disease states involving
CC (II). (II) is useful for generating antibodies against it, detecting or
CC quantitating a polypeptide in tissue, as molecular weight markers and as
CC a food supplement. (II) and its binding partners are useful in medical
CC imaging of sites expressing (II). (I) and (II) are useful for treating
CC disorders involving aberrant protein expression or biological activity.
CC The polypeptide and polynucleotide sequences have applications in
CC diagnostics, forensics, gene mapping, identification of mutations in
CC responsible for genetic disorders or other traits to assess biodiversity
CC and to produce other types of data and products dependent on DNA and
CC amino acid sequences. AAS6197-AAS94564 represent novel human
CC diagnostic coding sequences of the invention.
CC Note: The sequence data for this patent did not appear in the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/published_pct_sequences.
XX
XX SQ Sequence 1909 BP; 705 A; 443 C; 389 G; 372 T; 0 other;

Alignment Scores:
Pred. No.: 0.0134 Length: 1909
Score: 83.00 Matches: 18
Percent Similarity: 60.61% Conservative: 2
Best Local Similarity: 54.55% Mismatches: 13
Query Match: 43.01% Indels: 0
DB: 23 Gaps: 0

09-513-999c-7869_COPY_1_37 (1-37) x AAS88434 (1-1909)
OY 4 SerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuGlyProGluTyrVal 23
DB 205 TCAAGACCTTCCTGAGTACAGTCCATTGGCAGGACCGAGACGACGATACATA 146
OY 24 LysLeuGlyLeuGlyValCysLeuSerGlyCysSer 36
DB 145 AAGCCCTGGGTCTTGATGCTGCTGAGACAGCTGCTCC 107

RESULT 15
ID AAH98980 standard: cDNA, 609 BP.
XX
XX AAH98980;
AC
XX
XX 12-OCT-2001 (first entry)
DE Human EST-derived coding sequence SEQ ID NO: 837.
XX
XX Human: sheep; pig; cow; fruit fly; yeast; hamster; macaque; horse;
KW tomato; monkey; dog; sea urchin; expressed sequence tag; EST;
KW diagnostics; forensic test; gene mapping; genetic disorder;
KW biodiversity; gene therapy; nutrition; ss.
```

```
XX
XX Homo sapiens.
OS
XX
XX WO200154477-A2.
XX
XX 02-AUG-2001.
XX
XX 25-JAN-2001; 2001WO-US02687.
XX
XX 25-JAN-2000; 2000US-0491404.
PR 17-JUL-2000; 2000US-0617746.
PR 03-AUG-2000; 2000US-0631451.
PR 13-SEP-2000; 2000US-0663870.
XX
XX (HYSE-) HYSEQ INC.
XX
XX Tang YT, Liu C, Zhou P, Qian XB, Wang Z, Chen R, Asundi V;
PI Cao Y, Drmanac RA, Zhang J, Werhman T;
XX
XX WPI: 2001-476164/51.
DR P-PSDB: AAM24321.
XX
XX Isolated polypeptide for treatment of diseases, diagnostics, raising
PT antibodies and research use -
XX
XX Claim 1: Page 694; 1275pp; English.
XX
XX The present invention provides the protein and coding sequences of novel
CC proteins from a variety of organisms, including human, dog, cat, horse,
CC cow, pig, hamster, monkey, macaque, yeast, bacteria, fruit fly, sea
CC urchin and tomato. These were derived from expressed sequence tags (ESTs)
CC from the organism of interest. They can be used in diagnostics,
CC forensics, gene mapping, identification of mutations, to assess
CC biodiversity and for nutritional purposes. The present sequence is a cDNA
CC of the invention.
XX
XX SQ Sequence 609 BP; 99 A; 162 C; 173 G; 175 T; 0 other;

Alignment Scores:
Pred. No.: 0.0176 Length: 609
Score: 78.00 Matches: 17
Percent Similarity: 71.43% Conservative: 3
Best Local Similarity: 60.71% Mismatches: 8
Query Match: 40.41% Indels: 0
DB: 22 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AAH98980 (1-609)
OY 6 AlaLeuGlnAspSerPheSerLeuGlnGlyLeuGlyProGluTyrValLysLeu 25
DB 273 GCCTTGCCCTGAGATGCCATCACCTTCTTGGGATCCTAAGCTGTATGTAAAGCTC 332
OY 26 LeuGlyLeuGlyValCysLeuSer 33
DB 333 CTGGGTCTGTATGTGCTGAGC 356

Search completed: April 24, 2003, 23:01:50
Job time : 165 secs
```


GenCore version 5.1.4_p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - nucleic search, using frame_plus_p2n model

Run on: April 24, 2003, 22:34:13 ; Search time 1054 Seconds
(without alignments) 568.532 Million cell updates/sec

Title: US-09-513-999c-7869_COPY_1_37
Perfect score: 193
Sequence: 1 MGGSFALQDSRSSLQGLGPEYKVLGLCYCLSGCSTR 37

Scoring table: BLOSUM62
Xgapop 10.0 , Ygapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Indexed: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODE=frame+ p2n.model -DEV=xlh
-Q=ccp2.1/USPFC_pool/US09513999/unat_18042003.170937.28370/app_query.fasta.1.199
-DB=EST -QFMT=fastap -SUFFIX=p2n.rst -MINMATCH=0.1 -LOOPEXT=0 -LOOPEXT=0
-UNITS=bits -START=1 -END=1 -MATRIX=biosum62 -TRANS=human40.cdi -LIST=45
-DOCALIGN=200 -THR.SCORE=pct -THR.MAX=100 -THR.MIN=0 -ALIGN=15 -MODE=LOCAL
-OUTFMT=pct -NORM=ext -HEAPSIZE=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09513999 @CGN.1.1456 @unat.18042003.170937.28370 -NCPU=6 -ICPU=3
-NO_XLPXY -NO_MMAP -LARGQUERY -NEG.SCORES=0 -MATT -LONGLOG -DEV.TIMEOUT=120
-WARN.TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :
EST : *
1: em_estba : *
2: em_esthum : *
3: em_estlin : *
4: em_estlun : *
5: em_estlov : *
6: em_estlpl : *
7: em_estlro : *
8: em_hlcc : *
9: gb_estcl : *
10: gb_est2 : *
11: gb_hlc : *
12: gb_est3 : *
13: gb_est4 : *
14: gb_est5 : *
15: em_estfun : *
16: em_estom : *
17: gb_gss : *
18: em_gss_hum : *
19: em_gss_inv : *
20: em_gss_pln : *
21: em_gss_vrt : *
22: em_gss_fun : *
23: em_gss_mam : *
24: em_gss_mus : *
25: em_gss_other : *
26: em_gss_pro : *
27: em_gss_rtd : *

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
1	193	100.0	470	17	A0770688	A0770688 HS.5368_B
2	169	87.6	525	17	A0165256	A0165256 HS.3025_B
3	129	66.8	864	17	A0739814	A0739814 HS.3505_A
4	109	56.5	519	17	A0186743	A0186743 HS.3109_B
5	109	56.5	533	17	A0358845	A0358845 HS.5028_B
6	108	56.0	628	17	A0237815	A0237815 HS.11-70
7	105	54.4	292	17	A0508480	A0508480 RPT-11-2
8	105	54.4	632	17	AG146082	AG146082 Pan trogl
9	104	53.9	666	17	A0427698	A0427698 CITR1-E1
10	102	52.8	418	9	AA493535	AA493535 ng75g10.s
11	102	52.8	556	17	A0384817	A0384817 RPT11-13
12	102	52.8	695	17	AG179297	AG179297 Pan trogl
13	101	52.3	703	17	A0534396	A0534396 RPT-11-3
14	100	51.8	634	17	AG160901	AG160901 Pan trogl
15	100	51.8	659	17	AG151043	AG151043 Pan trogl
16	100	51.8	676	17	AG061401	AG061401 Pan trogl
17	100	51.8	769	17	A0899390	A0899390 HS.5234_A
18	99	51.3	362	17	A0102366	A0102366 HS.3040_A
19	99	51.3	410	17	A0442274	A0442274 HS.5137_A
20	99	51.3	416	17	A0182486	A0182486 HS.3077_A
21	99	51.3	453	17	A0437684	A0437684 HS.5137_A
22	99	51.3	485	17	A0671849	A0671849 HS.5462_A
23	99	51.3	635	17	A0390599	A0390599 CITR1-E1
24	98	50.8	546	17	A0435071	A0435071 HS.5114_B
25	98	50.3	597	17	A0506884	A0506884 RPT-11-1
26	97	50.0	410	17	A031865	A031865 RPT-11-3
27	96.5	50.0	672	17	A051272	A051272 RPT-11-4
28	96.5	49.7	347	17	A0631315	A0631315 RPT-11-4
29	96	49.7	444	17	A0463109	A0463109 HS.5211_A
30	96	49.7	663	17	AG091225	AG091225 Pan trogl
31	96	49.2	360	17	A0207172	A0207172 HS.3239_B
32	95	49.2	519	17	A0139984	A0139984 HS.5106_A
33	95	49.2	546	17	A0541696	A0541696 RPT-11-3
34	95	49.2	672	17	AG051939	AG051939 Pan trogl
35	95	49.2	363	17	A0120796	A0120796 HS.3076_A
36	94	48.7	425	17	A0683450	A0683450 HS.5432_B
37	94	48.7	452	17	A0534129	A0534129 RPT-11-3
38	94	48.7	516	17	A0457001	A0457001 HS.5151_A
39	94	48.7	546	17	A0333597	A0333597 HS.5008_A
40	94	48.2	529	17	A0881246	A0881246 HS.5137_B
41	93	47.7	580	17	A0532835	A0532835 RPT-11-3
42	92	47.2	463	17	A0550196	A0550196 RPT-11-4
43	91	47.2	546	17	A0468922	A0468922 HS.5139_A
44	91	47.2	559	17	A0385332	A0385332 RPT11-14
45						

ALIGNMENTS

RESULT 1
A0770688
LOCUS
DEFINITION HS.5368_B2.C08.SP6E RPT-11 Human Male BAC Library Homo sapiens
ACCESSION A0770688
VERSION A0770688.1 GI:5648804
KEYWORDS
SOURCE
ORGANISM
human.
human sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 470)
Mahatras,G.O., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T.,
Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and

JOURNAL
MEDLINE
COMMENT

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu

Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm) or from Research Genetics (info@resgen.com). BAC end Web Server: http://www.htsc.washington.edu

Plate: 944 row: F column: 16
Seq primer: SP6
Class: BAC ends
High quality sequence stop: 470.

FEATURES
source
location/Qualifiers
1..470
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=944 Col=16 Row=F"
/clone_lib="RPCI-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI; Male blood DNA was isolated from one randomly chosen donor and partially digested with a combination of EcoRI and EcoRI Methylase. Size selected DNA was cloned into the pBAC3.6 vector at EcoRI sites"

BASE COUNT 83 a 112 c 131 g 141 t 3 others

ORIGIN

Alignment Scores:
Pred. No.: 4,59e-16 Length: 470
Score: 193.00 Matches: 37
Percent Similarity: 100.00% Conservative: 0
Best Local Similarity: 100.00% Mismatches: 0
Query Match: 100.00% Indels: 0
Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AQ770688 (1-470)

Db 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyPro 20
|||||
Db 103 ATGGGTGATCTTTTGCCTTCAGAGATCTTTTCATCTTTGACAGGACCTTGGGGCGG 162
|||||

Qy 21 GltTyrValIysLeuGlnGlyLeuGlnCysValCysLeuSerGlyCysSerThr 37
|||||

Db 163 GAGTATGTAAACTCTGGTCTCTGTGTGTGCTGAGTGGCTCTCTACT 213
|||||

RESULT 2
LOCUS AQ165256 525 bp DNA linear GSS 16-JUL-1998
DEFINITION HS_3025_B2_G06_T7 CIT Approved Human Genomic Sperm Library D Homo sapiens genomic clone Plate=3025 Col=12 Row=N, DNA sequence.
ACCESSION AQ165256
VERSION AQ165256.1 GI:3563451
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 525)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

TITLE

JOURNAL
MEDLINE
COMMENT

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu

Sequence Tagged Connector
Plate: 3025 row: N column: 12
Class: BAC ends
High quality sequence stop: 525.

FEATURES
source
location/Qualifiers
1..525
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=3025 Col=12 Row=N"
/clone_lib="CIT Approved Human Genomic Sperm Library D"
/sex="male"
/note="Organ: sperm; Vector: pBeloBAC11; BAC clones in E-Coli DH10B"

BASE COUNT 102 a 139 c 137 g 143 t 4 others

ORIGIN

Alignment Scores:
Pred. No.: 8.15e-13 Length: 525
Score: 169.00 Matches: 33
Percent Similarity: 91.89% Conservative: 1
Best Local Similarity: 89.19% Mismatches: 3
Query Match: 87.56% Indels: 0
Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AQ165256 (1-525)

Qy 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyPro 20
|||||

Db 66 ATGGCGGATCTTCCCTTCAGGAATATTCATCTTTCAGAGGACCTTGGGGCGG 125
|||||

Qy 21 GltTyrValIysLeuGlnGlyLeuGlnCysValCysLeuSerGlyCysSerThr 37
|||||

Db 126 GAGTATGTAAACTCTGGTCTCTGTGTGTGCTGAGTGGCTCTCTACT 176
|||||

RESULT 3
LOCUS AQ739814 864 bp DNA linear GSS 16-JUL-1999
DEFINITION HS_3505_A1_A09_T7A RPCI-11 Human Male BAC library Homo sapiens genomic clone Plate=1081 Col=17 Row=A, DNA sequence.
ACCESSION AQ739814
VERSION AQ739814.1 GI:5517336
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
1 (bases 1 to 864)
Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome

TITLE

JOURNAL
MEDLINE
COMMENT

Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
Contact: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu

Clones are derived from the human BAC library RPCI-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from

BACPAC Resources (http://bacpac.med.buffalo.edu/ordering_bac.htm)
or from Research Genetics (<http://www.htsc.washington.edu>). BAC end Web Server:

Plate: 1081 row: A column: 17
Seq primer: T7

Class: BAC ends

High quality sequence stop: 864.

FEATURES
Location/Qualifiers
1. 864

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=1081 Col=17 Row=A"
/clone_lib="RPC1-11 Human Male BAC Library"
/sex="male"
/note="Vector: pBACE3.6; Site 1: EcoRI; Site 2: EcoRI;
Male blood DNA was isolated from one randomly chosen donor
and partially digested with a combination of EcoRI and
EcoRI Methylase. Size selected DNA was cloned into the
pBACE3.6 vector at EcoRI sites"

BASE COUNT 247 a 182 c 178 g 257 t

IN

Alignment Scores:

Prod. No.: 3.03e-07 Length: 864
Score: 129.00 Matches: 23
Percent Similarity: 92.86% Conservative: 3
Best Local Similarity: 82.14% Mismatches: 2
Query Match: 66.84% Indels: 0
DB: 17 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AQ739814 (1-864)

OY 9 AspSerPheSerLeuGlnGlyLeuGlyProGluTyrValLysLeuGlyLeu 28

DB 585 GACAAATTTTCGACCTGGAGGGGCTCGGGTCCACAGTATGTAACCTCGGGCCTC 526

OY 29 CysValCysLeuSerGlyCysSer 36

DB 525 TGGCTGTGCTGAGTGCTGCTCT 502

RESULT 4

LOCUS

DEFINITION HS_3109_B1.A06.T7 C1T Approved Human Genomic Sperm Library D Homo

ACCESION A0186743 sapiens genomic clone Plate=3109 Col=11 Row=B, DNA sequence.

VERSION A0186743.1 GI:3586185

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

CONTACT: Mahairas GG, Wallace JC, Hood L

High Throughput Sequencing Center

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Sequence Tagged Connector

Plate: 3109 row: B column: 11

Class: BAC ends

High quality sequence stop: 519.

Location/Qualifiers

1. 519

SOURCE

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="Plate=3109 Col=11 Row=B"
/clone_lib="C1T Approved Human Genomic Sperm Library D"
/sex="male"
/note="Organ: sperm; Vector: pBeloBAC11; BAC Clones in
E-Coli DH10B"

BASE COUNT 144 a 155 c 122 g 93 t 5 others

ORIGIN

Alignment Scores:

Prod. No.: 7.79e-05 Length: 519
Score: 109.00 Matches: 21
Percent Similarity: 82.76% Conservative: 3
Best Local Similarity: 72.41% Mismatches: 5
Query Match: 56.48% Indels: 0
DB: 17 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x A0186743 (1-519)

OY 8 GlnAspSerPheSerLeuGlnGlyLeuGlyProGluTyrValLysLeuGly 27

DB 399 AAGGATGGATCTTCTGCGCGGGAATCCTGGGCGTGAGATGTAACCTTGGGT 340

OY 28 LeuCysValCysLeuSerGlyCysSer 36

DB 339 CTCTGTGTGCTCCAGTGCTGCTCT 313

RESULT 5

LOCUS

DEFINITION HS_5028_B1.B02.T7 RPC111 Human Male BAC Library Homo sapiens

ACCESION A0358845 genomic clone Plate=604 Col=3 Row=D, DNA sequence.

VERSION A0358845.1 GI:4207721

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

MEDLINE

COMMENT

CONTACT: Mahairas GG, Wallace JC, Hood L

High Throughput Sequencing Center

University of Washington

401 Queen Anne Avenue North, Seattle, WA 98109, USA

Tel: (206) 616-3618

Fax: (206) 616-3887

Email: jwallace@u.washington.edu

Clones may be purchased from Research Genetics (<http://www.htsc.washington.edu>)

BAC end Web Server: <http://www.htsc.washington.edu>

Plate: 604 row: D column: 3

Seq primer: T7

Class: BAC ends

High quality sequence stop: 533.

Location/Qualifiers

1. 533

/organism="Homo sapiens"

/db_xref="taxon:9606"

/clone="Plate=604 Col=3 Row=D"

/clone_lib="RPC111 Human Male BAC Library"

/sex="male"

/cell_type="Lymphocytes"

/note="Vector: pBACE3.6; RPC111 Human Male BAC Library"

BASE COUNT 98 a 132 c 139 g 151 t 13 others

ORIGIN

Alignment Scores:

Pred. No.: 8.03e-05 Length: 533
Score: 109.00 Matches: 19
Percent Similarity: 82.14% Conservative: 4
Best Local Similarity: 67.86% Mismatches: 5
Query Match: 56.48% Indels: 0
DB: 17 Gaps: 0

US-09-513-999C-7869_COPY_1_37 (1-37) x AQ358845 (1-533)

Qy 9 AspergillusserleugllyleuenglyproglutryrVallylsleuenglyleu 28

Db 132 GATGATTCCACCTTGCTGGTATCTCTGGACACAGATTTTAACATCATGGTCTC 191

Qy 29 CysValCysLeuSerGlyCysSer 36

Db 192 TGTATGCTGCTGACAGATCTCT 215

RESULT 6
37815

DEFINITION AQ237815 628 bp DNA linear GSS 21-APR-1999
RPcII1-70H4.TK RPcI-11 Homo sapiens genomic clone RPcI-11-70H4, DNA
sequence.

ACCESSION AQ237815

VERSION AQ237815.1 GI:3670106

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

FEATURES

source

1. 628

/organism="Homo sapiens"

/db_xref="GDB:752667"

/db_xref="taxon:9606"

/clone="RPcI-11-70H4"

/clone_lib="RPcI-11"

/sex="Male"

/cell_type="Lymphocytes"

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;

RPcII1 Human Male BAC Library"

BASE COUNT 125 a 150 c 164 g 189 t

ORIGIN

Alignment Scores:

Pred. No.: 0.000132 Length: 628

Score: 108.00 Matches: 20

Percent Similarity: 91.67% Conservative: 2

Best Local Similarity: 83.33% Mismatches: 2

Query Match: 55.96% Indels: 0

DB: 17 Gaps: 0

US-09-513-999C-7869_COPY_1_37 (1-37) x AQ237815 (1-628)

Qy 14 LeuGlyGlyLeuGlyProglutryrVallylsleuenglyleuGlyCysValCysLeuSer 33

Db 136 TTGCGCCAGGCTCTGGGGCCAGATGTAAACATCTGGCTGTGTGTGCTGAGC 195

Qy 34 GlyCysSerThr 37

Db 196 GGCTGCTCTTCA 207

RESULT 7

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

COMMENT

JOURNAL

FEATURES

source

1. 292

/organism="Homo sapiens"

/db_xref="GDB:7612823"

/db_xref="taxon:9606"

/clone="RPcI-11-294M24"

/clone_lib="RPcI-11"

/sex="Male"

/cell_type="Lymphocytes"

/note="Vector: pBAC3.6; Site_1: EcoRI; Site_2: EcoRI;

RPcII1 Human Male BAC Library"

BASE COUNT 71 a 69 c 78 g 73 t

ORIGIN

Alignment Scores:

Pred. No.: 0.000138 Length: 292

Score: 105.00 Matches: 19

Percent Similarity: 91.67% Conservative: 3

Best Local Similarity: 79.17% Mismatches: 2

Query Match: 54.40% Indels: 0

DB: 17 Gaps: 0

US-09-513-999C-7869_COPY_1_37 (1-37) x AQ508480 (1-292)

Qy 13 SerLeuGlyLeuGlyProglutryrVallylsleuenglyleuGlyCysValCysLeu 32

Db 81 GCCTTCAGGGAGATCTGAGGCCAGATATCTAAACTCTGCTGTGTGTGCTGCTA 140

Qy 33 SerGlyCysSer 36

VERSION	AQ427698.1	GI:4500605
KEYWORDS	GSS.	
SOURCE	human.	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 666)	
TITLE	Zhao,S., Adams,M.D., Niernan,W., Malek,J., Shizuya,H., Simon,M. and Venter,J.C.	
JOURNAL	Use of BAC End Sequences from Caltech Libraries for Sequence-Ready Map Building	
COMMENT	Unpublished (1997) Other_GSSs: CTRBI-EI-2575E9.TF Contact: Shaying Zhao, William Niernan, Mark Adams Department of Eukaryotic Genomics The Institute for Genomic Research 9712 Medical Center Dr., Rockville, MD 20850 Tel: 301 838 0200 Fax: 301 838 0208 Email: hbe@tigr.org Clones are available from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/tdb/humgen/bac_end_search/bac_end_search.html . Seq primer: M13 Reverse Class: BAC ends.	
FEATURES	Location/Qualifiers	
Source	1..666 /organism="Homo sapiens" /db_xref="taxon:9606" /clone_id="2575E9" /clone_lib="CTRBI-EI" /sex="male" /contig="vector: pBelobAC11; site_1: EcoRI; site_2: EcoRI; Caltech Human BAC library D"	
BASE COUNT	136 a 157 c 171 g 201 t	1 others
ORIGIN		
Alignment Scores:	Pred. No.: 0.000479 Length: 666	
Score:	104.00 Matches: 22	
Percent Similarity:	76.47% Conservative: 4	
Best Local Similarity:	64.71% Mismatches: 8	
Query Match:	53.89% Indels: 0	
DB:	17 gaps: 0	
US-09-513-999C_COPY_1_37 (1-37) x AQ427698 (1-666)		
QY	3 G1ySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyProGluTyr 22	
Db	:::: : ::::: :::	
	365 GGACTTCACGCCCTCGCTGCAGATTGCAGACAAAGTTTGTTGGGATCCCTTTGGCCAGAGCTT 424	
QY	23 ValLysLeuLeuGlnGlyLeuCysValCysLeuSerG1yCysSer 36	
Db		
	425 GTAAGCTCCTCGGTCTGTGTGTGACTGACGAGCTGCTCT 466	
RESULT 10	AA493535/c	
LOCUS	AA493535 418 bp mRNA linear EST 18-AUG-1997	
DEFINITION	ng75g10.s1 NCI_CGAP_Pr6 Homo sapiens cDNA clone IMAGE:940674	
	similar to contains element PRT7 repetitive element ; , mRNA	
sequence.	AA493535	
ACCESSION	AA493535	
VERSION	AA493535.1	
KEYWORDS	EST	
SOURCE	human.	
ORGANISM	Homo sapiens	
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo. 1 (bases 1 to 418)	
TITLE	NCI-CGAP http://www.ncbi.nlm.nih.gov/cgi/gap . National Cancer Institute, Cancer Genome Anatomy Project (CGAP),	

PRIMERS
Sequencing: T7
LIBRARY
Vector : PBAC3.6
R.Site 1 : ECORI
R.Site 2 : EcoRI.

FEATURES

Source

1. 695
/organism="Pan troglodytes"
/db_xref="taxon:9598"
/clone_id="RP43-051111.T7"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RPCI-43 Chimpanzee Male BAC Library"
BASE COUNT 243 a 135 c 122 g 192 t 3 others
ORIGIN

Alignment Scores:

Pred. No.: 0.000929 Length: 695
Score: 102.00 Matches: 19
Percent Similarity: 78.57% Conservative: 3
Local Similarity: 67.86% Mismatches: 6
Query Match: 52.85% Indels: 0
DB: 17 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x AG179297 (1-695)

OY 9 AsperPheserSerLeuGlnGlyLeuLeuGlyProGluTyrValLysLeuGlyLeu 28

DB 135 GACGATTTCCCTCCGATCATGCGCCAGATATGTAAACCTCGGTCTC 194

OY 29 CysValCysLeuSerGlyCysSer 36

DB 195 TGTGTATCTGATGAGTGGCTGCTCT 218

RESULT 13

LOCUS A0534396 703 bp DNA linear GSS 18-MAY-1999

DEFINITION RPCI-11-38013.T7 RPCI-11 Homo sapiens genomic clone RPCI-11-38013,

ACCESSION A0534396

VERSION A0534396.1 GI:4846086

KEYWORDS GSS.

SOURCE human.

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.

1 (bases 1 to 703)

Zhao,S., Adams,M.D., Nierman,W., Malek,J., de Jong,P. and Venter

J.C.

Use of BAC End Sequences from Library RPCI-11 for Sequence-Ready

Map Building

Unpublished (1997)

Contact: Shaying Zhao, William Nierman, Mark Adams

Department of Eukaryotic Genomics

The Institute for Genomic Research

9712 Medical Center Dr., Rockville, MD 20850

Tel: 301 838 0200

Fax: 301 838 0208

Email: hbe@tigr.org

Clones are derived from the human BAC library RPCI-11. For BAC

library availability, please contact Pieter de Jong

(pieter@jeong.med.buffalo.edu). Clones may be purchased from

BACPAC Resources (http://bacpac.med.buffalo.edu/ordering) or from

Research Genet cs (info@resgen.com). BAC end search page:

http://www.tigr.org/tldb/humgen/bac_end_search/bac_end_search.html.

FEATURES

Source

1. 703
/organism="Homo sapiens"
/db_xref="GDB:7645730"
/db_xref="taxon:9606"

/clone="RPCI-11-38013"

/clone_lib="RPCI-11"

/sex="Male"

/cell_type="lymphocytes"

/note="Vector: PBAC3.6; Site_1: EcoRI; Site_2: EcoRI;

RPCI11 Human Male BAC Library"

BASE COUNT 201 a 194 c 165 g 143 t

ORIGIN

Alignment Scores:

Pred. No.: 0.00128 Length: 703
Score: 101.00 Matches: 17
Percent Similarity: 75.00% Conservative: 4
Best Local Similarity: 60.71% Mismatches: 7
Query Match: 52.33% Indels: 0
DB: 17 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x A0534396 (1-703)

OY 9 AsperPheserSerLeuGlnGlyLeuLeuGlyProGluTyrValLysLeuGlyLeu 28

DB 389 GATGATATCCCTCCGATCATGCGCCAGATATGTAAACCTCGGTCTC 330

OY 29 CysValCysLeuSerGlyCysSer 36

DB 329 TGTATGTCTGATGAGTGGCTGCTCT 306

RESULT 14

LOCUS AG160901 634 bp DNA linear GSS 09-JAN-2002

DEFINITION Pan troglodytes DNA, clone: RP43-026N23.T7, genomic survey

sequence.

ACCESSION AG160901

VERSION AG160901.1 GI:16690579

KEYWORDS GSS.

SOURCE Pan troglodytes

ORGANISM Pan troglodytes

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

Mammalia; Eutheria; Primates; Catarrhini; Homidae; Pan.

1

Fujiyama,A., Hattori,M., Toyoda,A., Taylor,T.D., Yada,T.,

Totoki,Y., Watanabe,H. and Sakaki,Y.

BAC end sequences of library RPCI-43

Unpublished

2 (bases 1 to 634)

Totoki,Y., Watanabe,H. and Sakaki,Y.

Direct Submission

Submitted (02-AUG-2001) Asao Fujiyama, The Institute of Physical

and Chemical Research (RIKEN), Genomic Sciences Center (GSC);

1-7-22 Suehiro-chou,Tsukumi-Ku, Yokohama, Kanagawa 230-0045, Japan

(E-mail:chimbos@resgen.com, riken-go.jp, URL:http://hgp.resgen.com/)

Tel:81-45-503-9111, Fax:81-45-503-9170

Clones are derived from the chimpanzee BAC library RPCI-43 This BAC

end was generated during the R&D process and may have higher chance

of clone tracking errors.

PRIMERS

Sequencing: T7

LIBRARY

Vector : PBAC3.6

R.Site 1 : EcoRI

FEATURES

Source

1. 634
/organism="Pan troglodytes"
/db_xref="taxon:9598"
/clone="RP43-026N23.T7"
/sex="male"
/cell_type="lymphocytes"
/clone_lib="RPCI-43 Chimpanzee Male BAC Library"
BASE COUNT 116 a 168 c 156 g 194 t
ORIGIN

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - nucleic search, using frame_plus.p2n model

Run on: April 24, 2003, 22:35:13 ; Search time 42 Seconds
(without alignments)
270.168 Million cell updates/sec

Title: US-09-513-999c-7869_COPY_1_37
Perfect score: 193
Sequence: 1 MGSEFALQDSFSIQILGILGPEYKVLGICVCLSGCST 37

Scoring table: BLOSUM62
Xgapop 10.0, Xgapext 0.5
Ygapop 10.0, Ygapext 0.5
Fgapop 6.0, Fgapext 7.0
Delop 6.0, Delext 7.0

atched: 441362 segs, 153338381 residues

Total number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODEL=frame.p2n.model -DEV=xlh
-O=/cg2n2.1/uspro_spool/US09513999/runat_18042003_170937_28379/app.query.fasta_1.199
-DB=Issued_Patents.NA -QFWT=fastap -SUFFIX=p2n.rni -MINMATCH=0.1 -LOOPL=0
-LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=blomsum62 -TRANS=human40.cdi
-LIST=45 -DOCALLIGN=200 -THR_SCORE=pct -THR_MAX=100 -THR_MIN=0 -ALIGN=15
-MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEAPSIZ=500 -MINLEN=0 -MAXLEN=2000000000
-USER=US09513999.ecgn1.1.32.ernat.18042003_170937_28379 -NCPU=6 -ICPU=3
-NO_XLPHY -NO_MAP -LARGEQUERY -NEG_SCORES=0 -WAIT -LONGLOG -DEV_TIMEOUT=120
-WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -Fgapop=6 -Fgapext=7
-YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database :

- 1: Issued_Patents.NA.*
- 2: /cg2n2.6/ptodata/1/ina/5A.COMB.seq.*
- 3: /cg2n2.6/ptodata/1/ina/5B.COMB.seq.*
- 4: /cg2n2.6/ptodata/1/ina/6A.COMB.seq.*
- 5: /cg2n2.6/ptodata/1/ina/6B.COMB.seq.*
- 6: /cg2n2.6/ptodata/1/ina/Backfiles1.seq.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length DB	ID	Description
1	58	30.1	2085	2	US-08-677-049-1
2	57	29.5	7379	4	US-09-341-587-5
3	56	29.0	2073	4	US-09-033-523-6
4	56	29.0	3120	1	US-08-456-647B-19
5	56	29.0	3120	2	US-08-456-647B-19
6	54.5	28.2	2064	3	US-08-875-944B-1
7	54.5	28.2	2064	4	US-09-116-049-3
8	54.5	28.2	2410	2	US-08-780-835B-1
9	54.5	28.2	2410	4	US-09-303-268-1
10	54.5	28.2	2410	4	US-09-116-049-1
11	54	28.0	16998	4	US-09-676-610B-24
12	53.5	27.7	1308	4	US-09-526-993-4

13	53.5	27.7	2167	4	US-09-526-993-3	Sequence 3, Appl1
14	53.5	27.7	2176	1	US-07-778-890A-2	Sequence 2, Appl1
15	53.5	27.7	3224	4	US-09-526-993-2	Sequence 2, Appl1
16	53.5	27.7	6070	4	US-09-526-993-1	Sequence 1, Appl1
17	53.5	27.7	6157	4	US-09-526-993-10	Sequence 10, Appl1
18	53.5	27.7	6202	4	US-09-526-993-8	Sequence 8, Appl1
19	52.5	27.2	59065	4	US-09-813-817-3	Sequence 3, Appl1
20	52.5	27.2	59065	4	US-09-813-817-3	Sequence 3, Appl1
21	52	26.9	2055	3	US-08-872-855-1	Sequence 3, Appl1
22	52	26.9	2800	3	US-08-872-855-1	Sequence 1, Appl1
23	51.5	26.7	1019	4	US-09-177-650-128	Sequence 128, App
24	51.5	26.7	1191	2	US-09-061-337-11	Sequence 11, Appl1
25	51.5	26.7	1191	2	US-09-122-129-11	Sequence 11, Appl1
26	51.5	26.7	1191	3	US-09-340-991-11	Sequence 11, Appl1
27	51.5	26.7	1191	3	US-08-974-609-11	Sequence 11, Appl1
28	51.5	26.7	1191	4	US-09-549-098-11	Sequence 11, Appl1
29	51.5	26.7	1297	3	US-09-083-521-4	Sequence 4, Appl1
30	51.5	26.7	3523	3	US-08-749-527-1	Sequence 4, Appl1
31	51	26.4	3390	1	US-08-453-742-26	Sequence 26, Appl1
32	51	26.4	3390	1	US-08-453-742-26	Sequence 26, Appl1
33	51	26.4	3390	1	US-08-453-742-26	Sequence 26, Appl1
34	51	26.4	3390	1	US-08-453-742-26	Sequence 26, Appl1
35	51	26.4	3390	1	US-08-452-802-26	Sequence 26, Appl1
36	51	26.4	3416	1	US-08-453-742-24	Sequence 24, Appl1
37	51	26.4	3416	1	US-08-454-664-24	Sequence 24, Appl1
38	51	26.4	3416	1	US-08-453-742-24	Sequence 24, Appl1
39	51	26.4	3416	1	US-08-452-802-24	Sequence 24, Appl1
40	51	26.4	3588	1	US-07-792-885A-2	Sequence 2, Appl1
41	51	26.4	9370	1	US-08-320-559-27	Sequence 27, Appl1
42	51	26.4	9370	5	US-08-345-860D-27	Sequence 27, Appl1
43	51	26.4	9391	1	PCT-US94-04496-27	Sequence 27, Appl1
44	51	26.4	9391	3	US-08-320-559-25	Sequence 25, Appl1
45	51	26.4	9391	5	US-08-545-860D-25	Sequence 25, Appl1

ALIGNMENTS

RESULT 1
US-08-677-049-1
Sequence 1, Application US/08677049
Patent No. 5858707
GENERAL INFORMATION:
APPLICANT: Guimaraes, M. Jorge
APPLICANT: Bazan, J. Fernando
APPLICANT: McClanahan, Terrill K.
TITLE OF INVENTION: ZICINIK, Albert
TITLE OF INVENTION: PURIFIED MAMMALIAN NUCLEOBASE PERMEASES;
NUMBER OF SEQUENCES: 12
CORRESPONDENCE ADDRESS:
ADDRESS: DNAX Research Institute
STREET: 901 California Avenue
CITY: Palo Alto
STATE: California
COUNTRY: USA
ZIP: 94304-1104
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/677,049
FILING DATE: 03-JUL-1996
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 60/000,788
FILING DATE: 03-JUL-1996
ATTORNEY/AGENT INFORMATION:
NAME: Ching, Edwin P.
REGISTRATION NUMBER: 34,090
REFERENCE/DOCKET NUMBER: DX0511
TELECOMMUNICATION INFORMATION:

Best Local Similarity: 51.52%

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-032-523-6 (1-2073)


```

: Patent No.6096542
:
: GENERAL INFORMATION:
: APPLICANT: FUJINAGA, Kei
: APPLICANT: YOSHIDA, Koichi
: APPLICANT: HIGASHINO, Fumihiro
: TITLE OF INVENTION: CANCER CONTROL
: NUMBER OF SEQUENCES: 5
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: BROWDY AND NEIMARK, P.L.L.C.
: STREET: 624 Ninth Street N.W., Ste. 300
: CITY: Washington
: STATE: D.C.
: COUNTRY: USA
: ZIP: 20001
:
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: PatentIn Release #1.0, Version #1.30
:
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/875,944B
: FILING DATE: 07-AUG-1997
: CLASSIFICATION: 514
:
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: JP 07-020173
: FILING DATE: 08-FEB-1995
:
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: PCT/JP96/00016
: FILING DATE: 09-JAN-1996
: ATTORNEY/AGENT INFORMATION:
: NAME: BROWDY, Roger L.
: REGISTRATION NUMBER: 25,618
: REFERENCE/DOCKET NUMBER: FUJINAGA-1
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (202) 628-5197
: TELEFAX: (202) 737-3528
:
: INFORMATION FOR SEQ ID NO: 1:
:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 2064 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
: MOLECULE TYPE: cDNA
: FEATURE:
: NAME/KEY: CDS
: LOCATION: 1..1386
: US-08-875-944B-1
:
: Alignment Scores:
:
: Seq. ID: 1
: Seq. ID: 2
: Seq. ID: 3
: Seq. ID: 4
: Seq. ID: 5
: Seq. ID: 6
: Seq. ID: 7
: Seq. ID: 8
: Seq. ID: 9
: Seq. ID: 10
: Seq. ID: 11
: Seq. ID: 12
: Seq. ID: 13
: Seq. ID: 14
: Seq. ID: 15
: Seq. ID: 16
: Seq. ID: 17
: Seq. ID: 18
: Seq. ID: 19
: Seq. ID: 20
: Seq. ID: 21
: Seq. ID: 22
: Seq. ID: 23
: Seq. ID: 24
: Seq. ID: 25
: Seq. ID: 26
: Seq. ID: 27
: Seq. ID: 28
: Seq. ID: 29
: Seq. ID: 30
: Seq. ID: 31
: Seq. ID: 32
: Seq. ID: 33
: Seq. ID: 34
: Seq. ID: 35
: Seq. ID: 36
: Seq. ID: 37
: Seq. ID: 38
: Seq. ID: 39
: Seq. ID: 40
: Seq. ID: 41
: Seq. ID: 42
: Seq. ID: 43
: Seq. ID: 44
: Seq. ID: 45
: Seq. ID: 46
: Seq. ID: 47
: Seq. ID: 48
: Seq. ID: 49
: Seq. ID: 50
: Seq. ID: 51
: Seq. ID: 52
: Seq. ID: 53
: Seq. ID: 54
: Seq. ID: 55
: Seq. ID: 56
: Seq. ID: 57
: Seq. ID: 58
: Seq. ID: 59
: Seq. ID: 60
: Seq. ID: 61
: Seq. ID: 62
: Seq. ID: 63
: Seq. ID: 64
: Seq. ID: 65
: Seq. ID: 66
: Seq. ID: 67
: Seq. ID: 68
: Seq. ID: 69
: Seq. ID: 70
: Seq. ID: 71
: Seq. ID: 72
: Seq. ID: 73
: Seq. ID: 74
: Seq. ID: 75
: Seq. ID: 76
: Seq. ID: 77
: Seq. ID: 78
: Seq. ID: 79
: Seq. ID: 80
: Seq. ID: 81
: Seq. ID: 82
: Seq. ID: 83
: Seq. ID: 84
: Seq. ID: 85
: Seq. ID: 86
: Seq. ID: 87
: Seq. ID: 88
: Seq. ID: 89
: Seq. ID: 90
: Seq. ID: 91
: Seq. ID: 92
: Seq. ID: 93
: Seq. ID: 94
: Seq. ID: 95
: Seq. ID: 96
: Seq. ID: 97
: Seq. ID: 98
: Seq. ID: 99
: Seq. ID: 100
: Seq. ID: 101
: Seq. ID: 102
: Seq. ID: 103
: Seq. ID: 104
: Seq. ID: 105
: Seq. ID: 106
: Seq. ID: 107
: Seq. ID: 108
: Seq. ID: 109
: Seq. ID: 110
: Seq. ID: 111
: Seq. ID: 112
: Seq. ID: 113
: Seq. ID: 114
: Seq. ID: 115
: Seq. ID: 116
: Seq. ID: 117
: Seq. ID: 118
: Seq. ID: 119
: Seq. ID: 120
: Seq. ID: 121
: Seq. ID: 122
: Seq. ID: 123
: Seq. ID: 124
: Seq. ID: 125
: Seq. ID: 126
: Seq. ID: 127
: Seq. ID: 128
: Seq. ID: 129
: Seq. ID: 130
: Seq. ID: 131
: Seq. ID: 132
: Seq. ID: 133
: Seq. ID: 134
: Seq. ID: 135
: Seq. ID: 136
: Seq. ID: 137
: Seq. ID: 138
: Seq. ID: 139
: Seq. ID: 140
: Seq. ID: 141
: Seq. ID: 142
: Seq. ID: 143
: Seq. ID: 144
: Seq. ID: 145
: Seq. ID: 146
: Seq. ID: 147
: Seq. ID: 148
: Seq. ID: 149
: Seq. ID: 150
: Seq. ID: 151
: Seq. ID: 152
: Seq. ID: 153
: Seq. ID: 154
: Seq. ID: 155
: Seq. ID: 156
: Seq. ID: 157
: Seq. ID: 158
: Seq. ID: 159
: Seq. ID: 160
: Seq. ID: 161
: Seq. ID: 162
: Seq. ID: 163
: Seq. ID: 164
: Seq. ID: 165
: Seq. ID: 166
: Seq. ID: 167
: Seq. ID: 168
: Seq. ID: 169
: Seq. ID: 170
: Seq. ID: 171
: Seq. ID: 172
: Seq. ID: 173
: Seq. ID: 174
: Seq. ID: 175
: Seq. ID: 176
: Seq. ID: 177
: Seq. ID: 178
: Seq. ID: 179
: Seq. ID: 180
: Seq. ID: 181
: Seq. ID: 182
: Seq. ID: 183
: Seq. ID: 184
: Seq. ID: 185
: Seq. ID: 186
: Seq. ID: 187
: Seq. ID: 188
: Seq. ID: 189
: Seq. ID: 190
: Seq. ID: 191
: Seq. ID: 192
: Seq. ID: 193
: Seq. ID: 194
: Seq. ID: 195
: Seq. ID: 196
: Seq. ID: 197
: Seq. ID: 198
: Seq. ID: 199
: Seq. ID: 200
: Seq. ID: 201
: Seq. ID: 202
: Seq. ID: 203
: Seq. ID: 204
: Seq. ID: 205
: Seq. ID: 206
: Seq. ID: 207
: Seq. ID: 208
: Seq. ID: 209
: Seq. ID: 210
: Seq. ID: 211
: Seq. ID: 212
: Seq. ID: 213
: Seq. ID: 214
: Seq. ID: 215
: Seq. ID: 216
: Seq. ID: 217
: Seq. ID: 218
: Seq. ID: 219
: Seq. ID: 220
: Seq. ID: 221
: Seq. ID: 222
: Seq. ID: 223
: Seq. ID: 224
: Seq. ID: 225
: Seq. ID: 226
: Seq. ID: 227
: Seq. ID: 228
: Seq. ID: 229
: Seq. ID: 230
: Seq. ID: 231
: Seq. ID: 232
: Seq. ID: 233
: Seq. ID: 234
: Seq. ID: 235
: Seq. ID: 236
: Seq. ID: 237
: Seq. ID: 238
: Seq. ID: 239
: Seq. ID: 240
: Seq. ID: 241
: Seq. ID: 242
: Seq. ID: 243
: Seq. ID: 244
: Seq. ID: 245
: Seq. ID: 246
: Seq. ID: 247
: Seq. ID: 248
: Seq. ID: 249
: Seq. ID: 250
: Seq. ID: 251
: Seq. ID: 252
: Seq. ID: 253
: Seq. ID: 254
: Seq. ID: 255
: Seq. ID: 256
: Seq. ID: 257
: Seq. ID: 258
: Seq. ID: 259
: Seq. ID: 260
: Seq. ID: 261
: Seq. ID: 262
: Seq. ID: 263
: Seq. ID: 264
: Seq. ID: 265
: Seq. ID: 266
: Seq. ID: 267
: Seq. ID: 268
: Seq. ID: 269
: Seq. ID: 270
: Seq. ID: 271
: Seq. ID: 272
: Seq. ID: 273
: Seq. ID: 274
: Seq. ID: 275
: Seq. ID: 276
: Seq. ID: 277
: Seq. ID: 278
: Seq. ID: 279
: Seq. ID: 280
: Seq. ID: 281
: Seq. ID: 282
: Seq. ID: 283
: Seq. ID: 284
: Seq. ID: 285
: Seq. ID: 286
: Seq. ID: 287
: Seq. ID: 288
: Seq. ID: 289
: Seq. ID: 290
: Seq. ID: 291
: Seq. ID: 292
: Seq. ID: 293
: Seq. ID: 294
: Seq. ID: 295
: Seq. ID: 296
: Seq. ID: 297
: Seq. ID: 298
: Seq. ID: 299
: Seq. ID: 300
: Seq. ID: 301
: Seq. ID: 302
: Seq. ID: 303
: Seq. ID: 304
: Seq. ID: 305
: Seq. ID: 306
: Seq. ID: 307
: Seq. ID: 308
: Seq. ID: 309
: Seq. ID: 310
: Seq. ID: 311
: Seq. ID: 312
: Seq. ID: 313
: Seq. ID: 314
: Seq. ID: 315
: Seq. ID: 316
: Seq. ID: 317
: Seq. ID: 318
: Seq. ID: 319
: Seq. ID: 320
: Seq. ID: 321
: Seq. ID: 322
: Seq. ID: 323
: Seq. ID: 324
: Seq. ID: 325
: Seq. ID: 326
: Seq. ID: 327
: Seq. ID: 328
: Seq. ID: 329
: Seq. ID: 330
: Seq. ID: 331
: Seq. ID: 332
: Seq. ID: 333
: Seq. ID: 334
: Seq. ID: 335
: Seq. ID: 336
: Seq. ID: 337
: Seq. ID: 338
: Seq. ID: 339
: Seq. ID: 340
: Seq. ID: 341
: Seq. ID: 342
: Seq. ID: 343
: Seq. ID: 344
: Seq. ID: 345
: Seq. ID: 346
: Seq. ID: 347
: Seq. ID: 348
: Seq. ID: 349
: Seq. ID: 350
: Seq. ID: 351
: Seq. ID: 352
: Seq. ID: 353
: Seq. ID: 354
: Seq. ID: 355
: Seq. ID: 356
: Seq. ID: 357
: Seq. ID: 358
: Seq. ID: 359
: Seq. ID: 360
: Seq. ID: 361
: Seq. ID: 362
: Seq. ID: 363
: Seq. ID: 364
: Seq. ID: 365
```

```

1 / Patent No.6248351
2 / GENERAL INFORMATION:
3 / APPLICANT: Hung, Men-Chie
4 / TITLE OF INVENTION: HUMAN PE3 IS A TUMOR SUPPRESSOR FOR CANCER CELLS
5 / FILE REFERENCE: UTSC:582
6 / CURRENT APPLICATION NUMBER: US/09/116,049A
7 / CURRENT FILING DATE: 1998-07-15
8 / NUMBER OF SEQ ID NOS: 11
9 / SOFTWARE: PatentIn Ver. 2.0
10 / SEQ ID NO 3
11 / LENGTH: 2064
12 / TYPE: DNA
13 / ORGANISM: Homo sapiens
14 / US-09-116-049-3
15
16 Alignment Scores:
17 Pred. No.: 66 Length: 2064
18 Score: 54.50 Matches: 15
19 Percent Similarity: 44.44% Conservative: 5
20 Best Local Similarity: 33.33% Mismatches: 12
21 Query Match: 28.24% Indels: 13
22 DB: 4 Gaps: 1
23
24 US-09-513-999C-7869-COPY_1_37 (1-37) x US-09-116-049-3 (1-2064)
25
26 QY 3 GlycerPhaeAlaLeuGlnAspSerPheSerLeuGlnGlyLeuGlyProGluTyr 22
27 Db 278 GCGCGCTTCGCGCTGCACAGACAGGCGCGCTGTGCGGAGACTCTGGCTCTTTG 219
28 QY 23 ValIysLeuLeuGlyLeu-----Cys 29
29 Db 218 ATCCGTGGTGGCGGCTGTGGAAGTACGTTTCTGAATGAAATCAGACAAACTGC 159
30 QY 30 ValCysLeuSerGly 34
31 Db 158 TCATCATCTGTCTGGT 144
32
33 RESULT 8
34 US-08-780-835B-1/c
35 / Sequence 1, Application US/08780835B
36 / Patent No.5922688
37 / GENERAL INFORMATION:
38 / APPLICANT: Hung, Men-Chie
39 / APPLICANT: Xing, Xiangming
40 / TITLE OF INVENTION: PE3 IS a Tumor Suppressor
41 / NUMBER OF SEQUENCES: 9
42 / CORRESPONDENCE ADDRESS:
43 / ADDRESSEE: ARNOLD, WHITE AND DURKEE
44 / STREET: P.O. Box 4433
45 / CITY: Houston
46 / STATE: Texas
47 / COUNTRY: USA
48 / ZIP: 77210-4433
49
50 / COMPUTER READABLE FORM:
51 / MEDIUM TYPE: Floppy disk
52 / COMPUTER: IBM PC compatible
53 / OPERATING SYSTEM: PC-DOS/MS-DOS
54 / SOFTWARE: PatentIn Release #1.0, Version #1.30
55 / CURRENT APPLICATION DATA:
56 / APPLICATION NUMBER: US/08/780,835B
57 / FILING DATE: 10-JAN-1997
58 / CLASSIFICATION: 514
59 / ATTORNEY/AGENT INFORMATION:
60 / NAME: Wilson, Mark B.
61 / REGISTRATION NUMBER: 37,259
62 / REFERENCE/DOCKET NUMBER: UTSC500
63 / TELECOMMUNICATION INFORMATION:
64 / TELEPHONE: (512) 418-3000
65 / TELEFAX: (512) 474-7577
66 / INFORMATION FOR SEQ ID NO: 1:
67 / SEQUENCE CHARACTERISTICS:
68 / LENGTH: 2410 base pairs
69 / type: nucleic acid
70

```



```

; STRANDEDNESS: single
; TOPOLOGY: linear
US-08-780-835B-1

Alignment Scores:
Pred. No.: 80.6 Length: 2410
Score: 54.50 Matches: 15
Percent Similarity: 44.44% Conservative: 5
Best Local Similarity: 33.33% Mismatches: 12
Query Match: 28.24% Indels: 13
DB: 2 Gaps: 1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-08-780-835B-1 (1-2410)

QY 3 GlySerPheAlaLeuGlnAspSerPheSerLeuGlnGlyLeuProGluTyr 22
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 694 GGTGGCTTCCTGCTGCACAGACAGGCGGCGGTCTGTGCGGGGACTGTGGTCTTGTG 635
      :: : :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: :: Cys 29
      634 ATTCGCTGGTGGGGGCTTATGAAAGCTAAGTTTCTGTAAGGAATCAGACAAACTGC 575

QY 30 ValCysLeuSergly 34
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 574 TCAATCAGCTGCCGT 560

RESULT 9
US-09-303-268-1/c
; Sequence 1, Application US/09303268
; Patent No. 6172212
; GENERAL INFORMATION:
; APPLICANT: Hung, Mien-Chie
; Xing, Xiangming
; TITLE OF INVENTION: PEA3 is a Tumor Suppressor
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: ARNOLD, WHITE AND DURKEE
; STREET: P.O. Box 4433
; CITY: Houston
; STATE: Texas
; COUNTRY: USA
; ZIP: 77210-4433
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/303,268
; FILING DATE: 30-Apr-1999
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/780,835
; FILING DATE: 10-JAN-1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Wilson, Mark B.
; REGISTRATION NUMBER: 37,259
; REFERENCE/DOCKET NUMBER: UTSC500
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (512) 418-3000
; TELEFAX: (512) 474-7577
; INFORMATION FOR SEQ ID NO: 1:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 2410 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-09-303-268-1

Alignment Scores:
Pred. No.: 80.6 Length: 2410
Score: 54.50 Matches: 15
Percent Similarity: 44.44% Conservative: 5
Best Local Similarity: 33.33% Mismatches: 12
Query Match: 28.24% Indels: 13
DB: 2 Gaps: 1

```

```

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-303-268-1 (1-2410)
Oy      3 GlycerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyProGluTyr 22
        ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db      694 GGTGGCTTCCTCGTCAGACAGAGCGGCGGTCTGTGCGGGACTCTGGGTTCCCTCTTG 635
Oy      23 ValLysLeuLeuGlyLeu-----Cys 29
        :: :|||||
Db      634 ATCCGTGGCGMGCGGCTATGAAGAAGCTAAGTTTCTGAAATGAATCAAGACAACAACCTGC 575
Oy      30 ValCysLeuSerGly 34
        ||||| |||||
Db      574 TCATCACCTGTCCGGT 560

RESULT 10
US-09-116-049-1/c
; Sequence 1, Application US/09116049A
; Patent No. 6248351
; GENERAL INFORMATION:
; APPLICANT: Hung, Men-Chie
; TITLE OF INVENTION: HUMAN PEAK IS A TUMOR SUPPRESSOR FOR CANCER CELLS
; FILE REFERENCE: UFGS:582
; CURRENT APPLICATION NUMBER: US/09/116, 049A
; CURRENT FILING DATE: 1998-07-15
; NUMBER OF SEQ ID NOS: 11
; SOFTWARE: Patentin Ver. 2.0
; SEQ ID NO 1
; LENGTH: 2410
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-116-049-1

Alignment Scores:
Pred. No.:          80.6           Length:       2410
Score:              54.50         Matches:      15
Percent Similarity: 44.44%        Conservative: 5
Best Local Similarity: 33.33%     Mismatches: 12
Query Match:        28.24%        Indels:    13
DB:                 4            gaps:        1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-116-049-1 (1-2410)
Oy      3 GlycerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyProGluTyr 22
        ||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db      694 GGTGGCTTCCTCGTCAGACAGAGCGGCGGTCTGTGCGGGACTCTGGGTTCCCTCTTG 635
Oy      23 ValLysLeuLeuGlyLeu-----Cys 29
        :: :|||||
Db      634 ATCCGTGGCGMGCGGCTATGAAGAAGCTAAGTTTCTGAAATGAATCAAGACAACAACCTGC 575
Oy      30 ValCysLeuSerGly 34
        ||||| |||||
Db      574 TCATCACCTGTCCGGT 560

RESULT 11
US-09-676-610B-24/c
; Sequence 24, Application US/09676610B
; Patent No. 6444465
; GENERAL INFORMATION:
; APPLICANT: C. Frank Bennett
; APPLICANT: Jacqueline Wyatt
; APPLICANT: Susan M. Freier
; TITLE OF INVENTION: OLIGONUCLEOTIDE INHIBITION OF HER-1 EXPRESSION
; FILE REFERENCE: RFS-0138
; CURRENT APPLICATION NUMBER: US/09/676, 610B
; CURRENT FILING DATE: 2000-09-29
; NUMBER OF SEQ ID NOS: 182
; SEQ ID NO 24
```

1	LENGTH: 169996
2	TYPE: DNA
3	ORGANISM: Homo sapiens
4	FEATURE:
5	NAME/KEY: exon
6	LOCATION: (1208)...(1472)
7	NAME/KEY: intron
8	LOCATION: (1473)...(124390)
9	NAME/KEY: exon
10	LOCATION: (124391)...(124544)
11	NAME/KEY: intron
12	LOCATION: (124545)...(125409)
13	NAME/KEY: exon
14	LOCATION: (125410)...(125595)
15	NAME/KEY: intron
16	LOCATION: (125596)...(128711)
17	NAME/KEY: exon
18	LOCATION: (128712)...(128848)
19	NAME/KEY: intron
20	LOCATION: (128849)...(133400)
21	NAME/KEY: exon
22	LOCATION: (133401)...(133469)
23	NAME/KEY: intron
24	LOCATION: (133470)...(134652)
25	NAME/KEY: exon
26	LOCATION: (134653)...(134773)
27	NAME/KEY: intron
28	LOCATION: (134774)...(136116)
29	NAME/KEY: exon
30	LOCATION: (136117)...(136261)
31	NAME/KEY: intron
32	LOCATION: (136262)...(137936)
33	NAME/KEY: exon
34	LOCATION: (137937)...(138053)
35	NAME/KEY: intron
36	LOCATION: (138054)...(138637)
37	NAME/KEY: exon
38	LOCATION: (138638)...(138766)
39	NAME/KEY: intron
40	LOCATION: (138767)...(138864)
41	NAME/KEY: exon
42	LOCATION: (138865)...(138940)
43	NAME/KEY: intron
44	LOCATION: (138941)...(139765)
45	NAME/KEY: exon
46	LOCATION: (139766)...(139860)
47	NAME/KEY: intron
48	LOCATION: (139861)...(142245)
49	NAME/KEY: exon
50	LOCATION: (142246)...(142445)
51	NAME/KEY: intron
52	LOCATION: (142446)...(143605)
53	NAME/KEY: exon
54	LOCATION: (143606)...(143738)
55	NAME/KEY: intron
56	LOCATION: (143739)...(145838)
57	NAME/KEY: exon
58	LOCATION: (145839)...(145931)
59	NAME/KEY: intron
60	LOCATION: (145932)...(147385)
61	NAME/KEY: exon
62	LOCATION: (147386)...(147544)
63	NAME/KEY: intron
64	LOCATION: (147545)...(153274)
65	NAME/KEY: exon
66	LOCATION: (153275)...(153321)
67	NAME/KEY: intron
68	LOCATION: (153322)...(155088)
69	NAME/KEY: exon
70	LOCATION: (155089)...(155231)
71	NAME/KEY: intron
72	LOCATION: (155232)...(156025)
73	NAME/KEY: exon

```

? LOCATION: (156026)...(156151)
? NAME/KEY: intron
? LOCATION: (156152)...(156826)
? NAME/KEY: exon
? LOCATION: (156927)...(156928)
? NAME/KEY: Intron
? LOCATION: (156929)...(163399)
? NAME/KEY: exon
? LOCATION: (163400)...(163586)
US-09-676-610B-24

Alignment Scores:
Pred. No.:      2.32e+04    Length:      169998
Score:          54.00       Matches:     12
Percent Similarity: 51.35%   Conservative: 7
Best Local Similarity: 32.43% Mismatches:    14
Query Match:      27.988     Indels:      4
DB:               4         Gaps:        1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-676-610B-24 (1-169998)
QY      3  GlySerPheAlauGlnAspSerPheSerSeriLeuIngLyLleuLeuGly----- 19
           ||| :|| |::| |::::| ::|||||::| |
Db  98748 GGATTCGTGCACCTGTCCTGATCGAGTGGCCACTTGCAAGAAGCATTAATTGGTTGTGATAT 98689
QY      20 --ProGUtRyValysLeuLeuGlyLeuCysValCysLeuSerGlyCys 35
            ||| ||| ||| :::|||||::| |
Db  98688 TGGCCAATAATTACAACCCTTGAAGTGTGTGTGTGTGTGWANGTGT 98638

RESULT 12
US-09-526-993-4/C
; Sequence 4, Application US/09526993
; Patent NO. 6465715
; GENERAL INFORMATION:
; APPLICANT: Zwaal, Richard
; APPLICANT: Aaseert, Wouter
; APPLICANT: Roelens, Ingele
; APPLICANT: Bogaert, Thierry
; TITLE OF INVENTION: EXPRESSION OF DNA OR PROTEINS IN C. ELEGANS
; FILE REFERENCE: B0192//012/ERC/KK
; CURRENT APPLICATION NUMBER: US/09/526,993
; EARLIEST FILING DATE: 2000-03-16
; EARLIER APPLICATION NUMBER: U.K. 9906018.8
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 4
; LENGTH: 1308
; TYPE: DNA
; ORGANISM: Caenorhabditis Elegans
US-09-526-993-4

Alignment Scores:
Pred. No.:      52              Length:      1308
Score:          53.50          Matches:     11
Percent Similarity: 64.29%     Conservative: 7
Best Local Similarity: 39.29%  Mismatches:    7
Query Match:      27.728      Indels:      3
DB:               4           Gaps:        1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-526-993-4 (1-1308)
QY      5  PheAlaLeuGlnAspSerPheSerSeriLeuIngLyLleuLeuGlyProGUtRyValys 24
           ||| |||::| ||||| |||::| |::::| ::|||::| |
Db  194 TTTCGATTTCAAACAGATTTCGTTAAATCGA-----ATTCGCCATATCATTCGC 144
QY      25 LeuLeuGlyLeuCysValCysValCys 32
           ||||| :::|| | ||||| 
Db  143 CTCCTGAAATATTTCTCGATGTTCG 120

RESULT 13
US-09-526-993-3/C
; Sequence 3, Application US/09526993
```

```

Patent No. 6465715
GENERAL INFORMATION:
APPLICANT: Zwaal, Richard
APPLICANT: Asaert, Moutier
APPLICANT: Roelens, Ingele
APPLICANT: Bogert, Thierly
TITLE OF INVENTION: EXPRESSION OF DNA OR PROTEINS IN C. ELEGANS
FILE REFERENCE: B0192/7012/ERG/KA
CURRENT APPLICATION NUMBER: US/09/526,993
CURRENT FILING DATE: 2000-03-16
EARLIER APPLICATION NUMBER: U.K. 9906018.8
EARLIER FILING DATE: 1999-03-16
NUMBER OF SEQ ID NOS: 11
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 3
LENGTH: 2167
TYPE: DNA
ORGANISM: Caenorhabditis Elegans
US-09-526-993-3
Alignment Scores:
Seq. No.:          99.8      Length:      2167
Score:             53.50     Matches:     11
Percent Similarity: 64.29%   Conservative: 7
Best Local Similarity: 39.29% Mismatches:      7
Query Match:       27.72%    Indels:      3
DB:                4        Gaps:          1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-526-993-3 (1-2167)
Oy      5  PhealaleuglnaaspserPheserSerleuglnGlyleuglnGlyProGluTyVallys 24
         |||:||||| |||:|||||
Db      194  TTTGATTTTCAAAACAGTTTCGTTTCATTCGAA-----ATTCCTCGATATCATTCGC 144
Oy      25  LeuleuglnGlyeCysValcysleu 32
         |||||:||||| |||||
Db      143  CTCCTTGATATTTGTCGATGTTTG 120

RESULT 14
US-07-778-890A-2/c
: Sequence 2, Application US/07778890A
: Patent No. 5266489
GENERAL INFORMATION:
APPLICANT: 1.REX-SENELONGE Arielle
APPLICANT: 2.KOHN Gillia
TITLE OF INVENTION: recombinant herpes viruses, in particular for the
TITLE OF INVENTION: production of vaccines, process for preparing them,
TITLE OF INVENTION: during this process and vaccines obtained.
NUMBER OF SEQUENCES: 7
CORRESPONDENCE ADDRESSES:
ADDRESSEE: Larson and Taylor
STREET: 727 Twenty-Third Street, South
CITY: Arlington
STATE: Virginia
COUNTRY: USA
ZIP: 22202
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette-5.25 inch, 360K storage
COMPUTER: IBM PC/XT/AT or compatibles
OPERATING SYSTEM: PC-DOS or MS-DOS version 2.2 or above
SOFTWARE: KEDIT or any ASCII Text Editor
CURRENT APPLICATION DATA: 07/778,890
APPLICATION NUMBER: US/07/778,890A
FILING DATE: 19920103
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: FR9003105, PCT/FR/91/00184
FILING DATE: 12-MAR-1990, 07-MAR-1991
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 2176 base pairs
TYPE: NUCLEIC ACID
STRANDEDNESS: double

```

Qy 25 LeuLeuGlyLeuCysValCysLeu 32
||||| : : |||
Db 1200 CTCCTGAAATATTGTCGATGTTG 1177

Search completed: April 24, 2003, 23:20:43
Job time : 58 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM protein - nucleic search, using frame_p2n model

Run on: April 24, 2003, 23:01:58 ; Search time 77 Seconds
(without alignments)
522.867 Million cell updates/sec

Title: US-09-513-999c-7869_COPY_1_37
Perfect score: 193
Sequence: 1 MGSEFALODSFSSIQGLGPRYKVLGICVCLSGCST 37

Scoring table:
BLOSUM62
Xgapop 10.0 , Xgapext 0.5
Ygapop 10.0 , Ygapext 0.5
Fgapop 6.0 , Fgapext 7.0
Delop 6.0 , Delext 7.0

Searched: 709820 seqs, 544064369 residues
Total number of hits satisfying chosen parameters: 1419640

Minimum DB seq Length: 0
Maximum DB seq Length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Command line parameters:
-MODEL=frame_p2n.model -DEV=xlh
-O=/cgn2_1/USPTO_SPOOL/US09513999/runat_18042003.170939.28423/app.query.fasta.1.199
-DB=Published_Applications_NA -OFMT=fastlap -SUFFIX=p2n.rnpb -MINMATCH=0.1
-LOOPEXT=0 -LOOPEXT=0 -UNITS=bits -START=1 -END=1 -MATRIX=bl0sum62
-TRANS=human40.cdi -LIST=45 -DOCALLIGN=200 -THR_SCORE=pct -THR_MAX=100
-THR_MIN=0 -ALIGN=15 -MODE=LOCAL -OUTFMT=ptc -NORM=ext -HEADSIZE=500 -MINLEN=0
-MAXLEN=2000000000 -USER=US09513999 @cgn 1.1.77 @runat.18042003.170939.28423
-NUPU=6 -ICPU=3 -NO_XLPXY -NO_MAP -LARGEUDERY -NEG_SCORES=0 -WAIT -LONKLOG
-DEV_TIMEOUT=120 -WARN_TIMEOUT=30 -THREADS=1 -XGAPOP=10 -XGAPEXT=0.5 -FGAPOP=6
-FGAPEXT=7 -YGAPOP=10 -YGAPEXT=0.5 -DELOP=6 -DELEXT=7

Database : Published Applications_NA:*

- 1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq:*
- 2: /cgn2_6/ptodata/1/pubpna/PCCT_NEW_PUB.seq:*
- 3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq:*
- 4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq:*
- 5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq:*
- 6: /cgn2_6/ptodata/1/pubpna/PCCTUS_PUBCOMB.seq:*
- 7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq:*
- 8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq:*
- 9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
- 10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq:*
- 11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*
- 12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq:*
- 13: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
- 14: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
C 1	93	48.2	684973	10 US-09-263-959-1	Sequence 1, Appli
C 2	88.5	45.9	570	10 US-09-864-761-9118	Sequence 9118, Ap
C 3	78	40.4	428	9 US-09-918-995-8153	Sequence 8153, Ap
C 4	69.5	36.0	412	10 US-09-783-590-1450	Sequence 1450, Ap

5	63	32.6	162	10 US-09-864-761-31393	Sequence 31393, A
6	63	31.6	519	10 US-09-864-761-14866	Sequence 14866, A
7	60.5	32.3	2730	10 US-09-965-528-27	Sequence 27, Appl
8	60.5	31.3	24533	9 US-09-764-868-1349	Sequence 1349, Ap
9	59.5	30.8	506	10 US-09-783-590-5391	Sequence 5391, Ap
C 10	57.5	29.8	1605	10 US-09-930-218-10	Sequence 10, Appl
11	57	29.5	1155	10 US-09-833-381-8	Sequence 8, Appl1
12	56	29.0	1380	10 US-09-974-300-685	Sequence 685, App
13	56	29.0	3120	10 US-09-158-722-19	Sequence 19, Appl
14	56	29.0	302250	10 US-09-962-832-154	Sequence 154, App
C 15	55	28.5	250	10 US-09-044-604-11	Sequence 11, Appl
C 16	55	28.5	360	10 US-09-867-701-8266	Sequence 8266, Ap
C 17	55	28.5	435	10 US-09-796-692-4049	Sequence 4049, Ap
18	55	28.5	467	10 US-09-864-761-6467	Sequence 6467, Ap
19	55	28.5	747	12 US-10-001-879-101	Sequence 101, App
C 20	55	28.5	855	10 US-09-044-604-11	Sequence 1, Appl1
21	55	28.5	3824	9 US-10-036-041-22	Sequence 22, Appl
22	55	28.5	3824	9 US-10-028-072-541	Sequence 541, App
23	55	28.5	3824	9 US-10-035-855-22	Sequence 22, Appl
24	55	28.5	3824	9 US-10-121-049-541	Sequence 541, App
25	55	28.5	3824	9 US-10-123-904-541	Sequence 541, App
26	55	28.5	3824	9 US-10-140-470-541	Sequence 541, App
27	55	28.5	3824	9 US-09-931-836-22	Sequence 22, Appl
28	55	28.5	3824	9 US-10-175-746-541	Sequence 541, App
29	55	28.5	3824	9 US-10-176-918-541	Sequence 541, App
30	55	28.5	3824	9 US-10-176-921-541	Sequence 541, App
31	55	28.5	3824	9 US-10-227-884-209	Sequence 209, App
32	55	28.5	3824	9 US-10-036-214-22	Sequence 22, Appl
33	55	28.5	3824	9 US-10-137-865-541	Sequence 541, App
34	55	28.5	3824	9 US-10-140-474-541	Sequence 541, App
35	55	28.5	3824	9 US-10-035-719-22	Sequence 22, Appl
36	55	28.5	3824	9 US-10-142-431-541	Sequence 541, App
37	55	28.5	3824	9 US-10-143-114-541	Sequence 541, App
38	55	28.5	3824	9 US-10-230-163-209	Sequence 209, App
39	55	28.5	3824	9 US-10-140-002-541	Sequence 541, App
40	55	28.5	3824	9 US-10-036-160-22	Sequence 22, Appl
41	55	28.5	3824	9 US-10-142-419-541	Sequence 541, App
42	55	28.5	3824	9 US-10-218-631-209	Sequence 209, App
43	55	28.5	3824	9 US-10-230-338-209	Sequence 209, App
44	55	28.5	3824	9 US-10-035-958-22	Sequence 22, Appl
45	55	28.5	3824	9 US-10-036-150-22	Sequence 22, Appl

ALIGNMENTS

RESULT 1
US-09-263-959-1/c
; Sequence 1, Application US/09263959
; Patent No. US20020150891A1
; GENERAL INFORMATION:
; APPLICANT: Hood, Leroy E.
; APPLICANT: Rowen, Lee
; TITLE OF INVENTION: DIAGNOSTIC AND THERAPEUTIC COMPOSITIONS AND METHODS WHICH U
; NUMBER OF SEQUENCES: 1279
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Seed and Berry LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: US
; ZIP: 98104-7092
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentin Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/263,959
; FILING DATE: 05-MAR-1999
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: McMasters, David D.

```

: REGISTRATION NUMBER: 33,963
: REFERENCE/DOCKET NUMBER: 920010.426C2
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (206) 622-4900
: TELEFAX: (206) 682-6031
: INFORMATION FOR SEQ ID NO: 1:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 684973 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: single
: TOPOLOGY: linear
:
: US-09-263-959-1

Alignment Scores:
Pred. No.: 0.0381 Length: 684973
Score: 93.00 Matches: 19
Percent Similarity: 75.00% Conservative: 2
Best Local Similarity: 67.86% Mismatches: 7
Query Match: 48.15% Indels: 0
Gaps: 10

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-263-959-1 (1-684973)
QY 9 AspSerPheSerSerLeuGlnGlyLeuGlyProGluTyrValIysLeuGlyLeu 28
Db 404731 GACAGATTCTGCTGCTGCTGGGATCTGAGGCTGGAATGTGTAACCTCTGGGTCTC 404672
QY 29 CysValCysLeuSerGlyCysSer 36
Db 404671 TGTGTGCTGTGAGCAGCTGCTCT 404648

RESULT 2
US-09-864-761-9118/C
: Sequence 9118, Application US/09864761
: Patent No. US20020048763A1
: GENERAL INFORMATION:
: APPLICANT: Penn, Sharon G.
: APPLICANT: Rank, David R.
: APPLICANT: Hanzel, David K.
: APPLICANT: Chen, Wensheng
: TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
: FILE REFERENCE: Aecmice-X-1
: CURRENT APPLICATION NUMBER: US/09/864,761
: PRIOR FILING DATE: 2001-05-23
: PRIOR APPLICATION NUMBER: US 60/180,312
: PRIOR FILING DATE: 2000-02-04
: PRIOR APPLICATION NUMBER: US 60/207,456
: PRIOR FILING DATE: 2000-05-26
: PRIOR APPLICATION NUMBER: US 09/632,366
: PRIOR FILING DATE: 2000-08-03
: PRIOR APPLICATION NUMBER: GB 24263.6
: PRIOR FILING DATE: 2000-10-04
: PRIOR APPLICATION NUMBER: US 60/236,359
: PRIOR FILING DATE: 2000-09-27
: PRIOR APPLICATION NUMBER: PCT/US01/00666
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00667
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00664
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00669
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00665
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00668
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00663
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00662
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00661
: PRIOR FILING DATE: 2001-01-30
```

```

: PRIOR APPLICATION NUMBER: PCT/US01/00670
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: US 60/234,687
: PRIOR FILING DATE: 2000-09-21
: PRIOR APPLICATION NUMBER: US 09/608,408
: PRIOR FILING DATE: 2000-06-30
: PRIOR APPLICATION NUMBER: US 09/774,203
: PRIOR FILING DATE: 2001-01-29
: NUMBER OF SEQ ID NOS: 49117
: SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
: SEQ ID NO 9118
: LENGTH: 570
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: OTHER INFORMATION: MAP TO AP000053.1
: OTHER INFORMATION: EXPRESSED IN HELA, SIGNAL = 2.8
: OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 2.4
: OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 4.8
: OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 4
: OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 2.3
: OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 3
: OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 3.6
: OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 4.4
:
: US-09-864-761-9118

Alignment Scores:
Pred. No.: 2.61e-05 Length: 570
Score: 88.50 Matches: 21
Percent Similarity: 61.11% Conservative: 1
Best Local Similarity: 58.33% Mismatches: 7
Query Match: 45.85% Indels: 7
Gaps: 10

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-864-761-9118 (1-570)
QY 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyPro 20
Db 568 ATGAATGATCT-----CTTCCTTGCTGGGATTCCTTGAGCTG 530
QY 21 GluTyrValIysLeuGlyLeuGlyCysValCysLeuSerGlyCysSer 36
Db 529 GAGTATGTAAATTCCTGCGGTCTTGTGTGCTGTGAGTGGCGGCTCT 482

RESULT 3
US-09-918-995-8153
: Sequence 8153, Application US/09918995
: Publication No. US20030073623A1
: GENERAL INFORMATION:
: APPLICANT: Hyseq, Inc.
: TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
: FILE REFERENCE: 20411-756
: CURRENT APPLICATION NUMBER: US/09/918,995
: PRIOR FILING DATE: 2001-07-30
: PRIOR APPLICATION NUMBER: US/09/235,076
: PRIOR FILING DATE: 1999-01-20
: NUMBER OF SEQ ID NOS: 38054
: SOFTWARE: FastSeq for Windows Version 3.0
: SEQ ID NO 8153
: LENGTH: 428
: TYPE: DNA
: ORGANISM: Homo sapiens
:
: US-09-918-995-8153

Alignment Scores:
Pred. No.: 0.000899 Length: 428
Score: 78.00 Matches: 17
Percent Similarity: 71.43% Conservative: 3
Best Local Similarity: 60.71% Mismatches: 8
Query Match: 40.41% Indels: 0
Gaps: 9
```

```

US-09-513-9996-7869_COPY_1_37 (1-37) x US-09-918-995-8153 (1-428)
OY 6 A1aLeuclnaspserPheSerSerLeuGlnGlyLeuGlyProGluTyrValLysLeu 25
Db 273 GCCTTGCCATGATGCCATCACCTTCTTGTGGGATCTTAAGCTGTGAATGAAGCTC 332
OY 26 LeuGlyLeuGlyValCysLeuSer 33
Db 333 CTGGGCTCTGTATGTGCTGAGC 356

RESULT 4
US-09-783-590-1450
; Sequence 1450, Application US/09783590
; Patent No. US20020110850A1
; GENERAL INFORMATION:
; APPLICANT: Dillon, Patrick J.
; APPLICANT: Haseltine, William A.
; APPLICANT: Li, Haodong
; APPLICANT: Rosen, Craig A.
; APPLICANT: Ruben, Steven M.
; TITLE OF INVENTION: Human Genes, Sequences, and Expression Products 16.2
; FILE REFERENCE: PO-15,2c1
; CURRENT APPLICATION NUMBER: US/09/783,590
; CURRENT FILING DATE: 2000-02-15
; PRIOR APPLICATION NUMBER: 08/420,856
; PRIOR FILING DATE: 1995-04-12
; PRIOR APPLICATION NUMBER: 08/346,731
; PRIOR FILING DATE: 1994-11-21
; NUMBER OF SEQ ID NOS: 12485
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1450
; LENGTH: 412
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (3)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (189)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (190)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (231)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (265)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (266)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (268)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (269)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (274)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (275)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (339)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (340)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc feature
; LOCATION: (370)

```

```

: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc feature
: LOCATION: (399)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc feature
: LOCATION: (402)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc feature
: LOCATION: (403)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc feature
: LOCATION: (406)
: OTHER INFORMATION: n equals a,t,g, or c
US-09-783-590-1450

Alignment Scores:
Pred. No.: 0.0202 Length: 412
Percent: 69.50 Matches: 16
Score Similarity: 61.29% Conservative: 3
Best Local Similarity: 51.61% Mismatches: 7
Query Match: 36.01% Indels: 5
DB: 10 Gaps: 1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-783-590-1450 (1-412)
Oy 11 PheserSerLeuGln-----c-GlyLeuGluGlyProGluTyValIysLeu 25
Db 172 TTTGCTGAGTTGCACCTNNTTTTTCTGTGGGAGCCCTGGAAAGCCAGAGATATCTGAGCCTN 231
Oy 26 LeuGlyLeuGysValCysLeuSerGlyCysSer 36
Db 232 TTGAATCTGTGCACAGGCGCTCAGTGCGTCCT 264

RESULT 5
US-09-864-761-31393
: Sequence 31393, Application US/09864761
: Patent No. US2002048763A1
: GENERAL INFORMATION:
: APPLICANT: Penn, Sharon G.
: APPLICANT: Rank, David R.
: APPLICANT: Hanzel, David K.
: APPLICANT: Chen, Wensheng
: TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
: FILE REFERENCE: Aecmica-x-1
: CURRENT APPLICATION NUMBER: US/09/864,761
: CURRENT FILING DATE: 2001-05-23
: PRIOR APPLICATION NUMBER: US 60/180,312
: PRIOR FILING DATE: 2000-02-04
: PRIOR APPLICATION NUMBER: US 60/207,456
: PRIOR FILING DATE: 2000-05-26
: PRIOR APPLICATION NUMBER: US 09/632,366
: PRIOR FILING DATE: 2000-08-03
: PRIOR APPLICATION NUMBER: GB 24263.6
: PRIOR FILING DATE: 2000-10-04
: PRIOR APPLICATION NUMBER: US 60/236,359
: PRIOR FILING DATE: 2000-09-27
: PRIOR APPLICATION NUMBER: PCT/US01/00666
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00667
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00664
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00669
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00665
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00668
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00663
: PRIOR FILING DATE: 2001-01-30
: PRIOR APPLICATION NUMBER: PCT/US01/00662
: PRIOR FILING DATE: 2001-01-30

```

```

; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 31393
; LENGTH: 162
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AC017089.2
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.8
; OTHER INFORMATION: SWISSPROT HIT: P98161, EVALUE 3.00e+00
; OTHER INFORMATION: EST_HUMAN HIT: A1792950.1, EVALUE 6.00e-07
; OTHER INFORMATION: NT HIT: AL163210.2, EVALUE 4.00e-04
; US-09-864-761-31393

Alignment Scores:
Pred. No.: 0.0695 Length: 162
Score: 63.00 Matches: 13
Percent Similarity: 66.67% Conservative: 1
Best Local Similarity: 61.90% Mismatches: 7
Query Match: 32.64% Indels: 0
DB: 10 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-864-761-31393 (1-162)
Qy 16 G1yLeuLeuG1yProG1yTyrVal1yLysLeuLeuG1yLeuCy5ValCy5LeuSerG1yCy5 35
    ||| |||||||||:||||| ||||||||| |||
Db 60 GGAAGTCCAGGCGCAGATATCTTAAGCTTTCATGTGTGCTGAGCGAGATGC 119

Qy 36 Ser 36
    |||
Db 120 TCT 122

RESULT 6
US-09-864-761-14866
; Sequence 14866, Application US/09864761
; Patent No. US20020048763A1
; GENERAL INFORMATION:
; APPLICANT: Penn, Sharon G.
; APPLICANT: Rank, David R.
; APPLICANT: Hanzel, David K.
; APPLICANT: Chen, Wensheng
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
; FILE REFERENCE: Aecmica-X-1
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
; PRIOR APPLICATION NUMBER: US 60/180,312
; PRIOR FILING DATE: 2000-02-04
; PRIOR APPLICATION NUMBER: US 60/207,456
; PRIOR FILING DATE: 2000-05-26
; PRIOR APPLICATION NUMBER: US 09/632,366
; PRIOR FILING DATE: 2000-08-03
; PRIOR APPLICATION NUMBER: GB 24263.6
; PRIOR FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: US 60/236,359
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: PCT/US01/00666
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00667
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00664
; PRIOR FILING DATE: 2001-01-30
```

```

; PRIOR APPLICATION NUMBER: PCT/US01/00669
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00665
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00668
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00663
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00662
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00661
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: PCT/US01/00670
; PRIOR FILING DATE: 2001-01-30
; PRIOR APPLICATION NUMBER: US 60/234,687
; PRIOR FILING DATE: 2000-09-21
; PRIOR APPLICATION NUMBER: US 09/608,408
; PRIOR FILING DATE: 2000-06-30
; PRIOR APPLICATION NUMBER: US 09/774,203
; PRIOR FILING DATE: 2001-01-29
; NUMBER OF SEQ ID NOS: 49117
; SOFTWARE: Annomax Sequence Listing Engine vers. 1.1
; SEQ ID NO 14866
; LENGTH: 519
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; OTHER INFORMATION: MAP TO AC017089.2
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.8
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
; US-09-864-761-14866

Alignment Scores:
Pred. No.: 0.303 Length: 519
Score: 63.00 Matches: 13
Percent Similarity: 66.67% Conservative: 1
Best Local Similarity: 61.90% Mismatches: 7
Query Match: 32.64% Indels: 0
DB: 10 Gaps: 0

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-864-761-14866 (1-519)
Qy 16 G1yLeuLeuG1yProG1yTyrVal1yLysLeuLeuG1yLeuCy5ValCy5LeuSerG1yCy5 35
    ||| |||||||||:||||| ||||||||| |||
Db 309 GGAAGTCCAGGCGCAGATATCTTAAGCTTTCATGTGTGCTGAGCGAGATGC 368

Qy 36 Ser 36
    |||
Db 369 TCT 371

RESULT 7
US-09-965-528-27
; Sequence 27, Application US/09965528
; Publication No. US20020187523A1
; GENERAL INFORMATION:
; APPLICANT: INCYTE GENOMICS, INC.
; APPLICANT: TANG, Y. Tom
; APPLICANT: YUE, Henry
; APPLICANT: LAL, Preeti
; APPLICANT: BURFORD, Neil
; APPLICANT: BANDMAN, Olga
; APPLICANT: BAUMZAI, Mariah R.
; APPLICANT: AZIMZAI, Yalda
; APPLICANT: LU, Dying Alpha M.
; APPLICANT: PATTERSON, Chandra
; TITLE OF INVENTION: EXTRACELLULAR SIGNALING MOLECULES
; FILE REFERENCE: PF-0701 USA
; CURRENT APPLICATION NUMBER: US/09/965,528
; CURRENT FILING DATE: 2001-09-26
; PRIOR APPLICATION NUMBER: 60/134,949
; PRIOR FILING DATE: 1999-05-19
; PRIOR APPLICATION NUMBER: 60/144,270
; PRIOR FILING DATE: 1999-07-15
```



```

; PRIOR APPLICATION NUMBER: 60/146,700
; PRIOR FILING DATE: 1999-07-30
; PRIOR APPLICATION NUMBER: 60/157,508
; PRIOR FILING DATE: 1999-10-04
; NUMBER OF SEQ ID NOS: 55
; SOFTWARE: PERL Program
; SEQ ID NO 27
; LENGTH: 2730
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; OTHER INFORMATION: Incybe ID No. US20020187523A1 1288847CB1
US-09-965-528-27

Alignment Scores:
Pred. No.: 6.24 Length: 2730
Score: 60.50 Matches: 13
Percent Similarity: 59.26% Conservative: 3
Best Local Similarity: 48.15% Mismatches: 10
Query Match: 31.35% Indels: 1
Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-965-528-27 (1-2730)
Qy 10 SerPheSerLeuGlnGlyLeuLeu--GlyProGluTyValysLeuLeuGlyLeu 28
Db 630 ACCTTCCTAGAGGCTCTCAGGGGCTCTCTCAGTCCCATCTCATGAGAACTAGTGGTTC 689

Qy 29 CysValCysLeuSerGlyCys 35
Db 690 TCCCTGATGACCAAGGGGTGT 710

RESULT 8
US-09-764-868-1349
; Sequence 1349, Application US/09764868
; Patent No. US2002016871A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT32
; CURRENT APPLICATION NUMBER: US/09/764,868
; CURRENT FILING DATE: 2001-01-17
; Prior application data removed - refer to PAM or file wrapper
; NUMBER OF SEQ ID NOS: 1510
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 1349
; LENGTH: 24533
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-868-1349

Alignment Scores:
Pred. No.: 100 Length: 24533
Score: 60.50 Matches: 16
Percent Similarity: 60.00% Conservative: 5
Best Local Similarity: 45.71% Mismatches: 13
Query Match: 31.35% Indels: 1
Gaps: 1

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-764-868-1349 (1-24533)
Qy 1 MetGlyGlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyPro 20
Db 14224 CTGGGGTGGGCTGTGGCCCTTCTAGCGCTCGTGGTGGGCCCAAGAGCTTTGGGTCT 14283

Qy 21 GluTyValysLeuLeuGlnGlyLeuGlyCysValCysLeuSerGlyCys 35
Db 14284 GAG--ATGCAACTGCTTGAGCTGAGCCGCGGATGATGATGAGCTGC 14325

RESULT 9
US-09-783-590-5991
; Sequence 5991, Application US/09783590

```

```

; Patent No. US20020110850A1
; GENERAL INFORMATION:
; APPLICANT: Dillon, Patrick J.
; APPLICANT: Haseltine, William A.
; APPLICANT: Li, Haodong
; APPLICANT: Rosen, Craig A.
; APPLICANT: Ruben, Steven M.
; TITLE OF INVENTION: Human Genes, Sequences, and Expression Products 16.2
; FILE REFERENCE: PO-16,201
; CURRENT APPLICATION NUMBER: US/09/783,590
; CURRENT FILING DATE: 2000-02-15
; PRIOR APPLICATION NUMBER: 08/420,856
; PRIOR FILING DATE: 1995-04-12
; PRIOR APPLICATION NUMBER: 08/346,731
; PRIOR FILING DATE: 1994-11-21
; NUMBER OF SEQ ID NOS: 12485
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 5991
; LENGTH: 506
; TYPE: DNA
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc_feature
; LOCATION: (110)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (331)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (351)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (364)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (402)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (435)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (460)
; OTHER INFORMATION: n equals a,t,g, or c
; NAME/KEY: misc_feature
; LOCATION: (502)
; OTHER INFORMATION: n equals a,t,g, or c
US-09-783-590-5991

Alignment Scores:
Pred. No.: 1.08 Length: 506
Score: 59.50 Matches: 16
Percent Similarity: 55.26% Conservative: 5
Best Local Similarity: 42.11% Mismatches: 8
Query Match: 30.83% Indels: 9
Gaps: 2

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-783-590-5991 (1-506)
Qy 3 GlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeu----- 18
Db 134 GGAGAAAAAGGACGACGATTAAGTTCGCGTGGCCCTTCACCAAGTGTGG 193

Qy 19 -----GlyProGluTyValysLeuLeuGlnGlyLeuGlyCysValCysLeu 32
Db 194 GAATTCTACTAGGTGAGAGATAC---AAATTAATTGAGAGTTTCATCTGTTTG 244

RESULT 10
US-09-930-218-10/c
; Sequence 10, Application US/09930218
; Patent No. US20020034810A1
; GENERAL INFORMATION:
; APPLICANT: goldsmith, orit
; APPLICANT: pecker, iris

```

```
APPLICANT: vlodavsky, israel
APPLICANT: israel, michal
TITLE OF INVENTION: AVIAN AND REPTILE DERIVED POLYNUCLEOTIDE ENCODING A POLYPEPTIDE
TITLE OF INVENTION: HEPARANASE ACTIVITY
FILE REFERENCE: 01/22335
CURRENT APPLICATION NUMBER: US/09/930,218
CURRENT FILING DATE: 2001-08-16
PRIOR APPLICATION NUMBER: 09/666,390
PRIOR FILING DATE: 2000-09-20
NUMBER OF SEQ ID NOS: 16
SOFTWARE: PatentIn version 3.1
SEQ ID NO 10
LENGTH: 1605
TYPE: DNA
ORGANISM: Gallus gallus
US-09-930-218-10
```

```
Alignment Scores:
Pred. No.: 9.73 Length: 1605
Score: 57.50 Matches: 11
Percent Similarity: 73.68% Conservative: 3
Best Local Similarity: 57.89% Mismatches: 4
Query Match: 29.79% Indels: 1
DB: 10 Gaps: 1
```

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-930-218-10 (1-1605)

```
OY 20 ProglutryValylsLeuLeuGlyLeuGlyCys---ValcysLeuSerglyCysSerThr 37
|||||  |||||  |||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 819 CCAAGATTTCAGAGAGCTTGAGCAGGCTGCTGGCTGGCTGGCGGGCTGCCCCACG 763
```

```
RESULT 11
US-09-833-381-8
; Sequence 8, Application US/09833381
; Patent No. US20020132090A1
; GENERAL INFORMATION:
; APPLICANT: Robison, Keith E.
; FILE OF INVENTION: No. US20020132090A1el Nucleic Acid and Protein Homologs
; TITLE REFERENCE: 5800-119
; CURRENT APPLICATION NUMBER: US/09/833,381
; CURRENT FILING DATE: 2001-04-11
; PRIOR APPLICATION NUMBER: 09/516,448
; PRIOR FILING DATE: 2000-02-29
; NUMBER OF SEQ ID NOS: 2050
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8
; LENGTH: 1155
; TYPE: DNA
; ORGANISM: Homo sapiens
09-833-381-8
```

```
Alignment Scores:
Pred. No.: 7.73 Length: 1155
Score: 57.00 Matches: 12
Percent Similarity: 53.66% Conservative: 10
Best Local Similarity: 29.27% Mismatches: 13
Query Match: 29.53% Indels: 6
DB: 10 Gaps: 1
```

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-833-381-8 (1-1155)

```
OY 1 MetcIyGlySerPheAlaLeuGlnAspSerPheSerSerLeu----- 14
:::|||||:::  |||  ::  ::::|
DB 535 GTAGAGAGCACTATACCTTGATCATGCCAATAATGCCCTCGCTCGAGCTCTGCAG 594
OY 15 GlngIyLeuLeuGlyProGluTrValylsLeuLeuGlyLeuGlyValcysLeuSergly 34
|||  ::  |||  ::  |||  |||||  |||||  |||||  |||||  |||||
DB 595 CAAAGCGCTACAGTATCCCTCGTACATGAGGCCCTTGCCCTGTGCATCTGTGGCGCTG 654
OY 35 Cys 35
|||
DB 655 TGT 657
```

```
RESULT 12
US-09-974-300-685
; Sequence 685, Application US/09974300
; Patent No. US20020146721A1
; GENERAL INFORMATION:
; APPLICANT: Berka, Randy M.
; APPLICANT: Clausen, Id Groth
; TITLE OF INVENTION: Methods For Monitoring Multiple Gene
; TITLE OF INVENTION: Expression
; FILE REFERENCE: 10085,500-US
; CURRENT APPLICATION NUMBER: US/09/974,300
; CURRENT FILING DATE: 2001-10-05
; PRIOR APPLICATION NUMBER: 09/680,598
; PRIOR FILING DATE: 2000-10-06
; PRIOR APPLICATION NUMBER: 60/279,526
; PRIOR FILING DATE: 2001-03-27
; NUMBER OF SEQ ID NOS: 8481
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 685
; LENGTH: 1380
; TYPE: DNA
; ORGANISM: Bacillus licheniformis
US-09-974-300-685
```

```
Alignment Scores:
Pred. No.: 14 Length: 1380
Score: 56.00 Matches: 9
Percent Similarity: 75.00% Conservative: 3
Best Local Similarity: 56.25% Mismatches: 4
Query Match: 29.02% Indels: 0
DB: 10 Gaps: 0
```

US-09-513-999c-7869_COPY_1_37 (1-37) x US-09-974-300-685 (1-1380)

```
OY 20 ProglutryValylsLeuLeuGlyLeuGlyCysValcysLeuSerglyCys 35
|||||  |||||  |||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
DB 564 CCGAGACTTGTCATCTTGAGGAGCTTGCACTGCATCTGCATCAGCGAGTGT 611
```

```
RESULT 13
US-09-158-722-19
; Sequence 19, Application US/09158722
; Publication No. US20030013848A1
; GENERAL INFORMATION:
; APPLICANT: Lemke Ph.D. et al., Greg E.
; TITLE OF INVENTION: PROTEIN-TYROSINE KINASE GENES
; NUMBER OF SEQUENCES: 54
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson P.C.
; STREET: 4225 Executive Square, Suite 1400
; CITY: La Jolla
; STATE: CA
; COUNTRY: US
; ZIP: 92037
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/158,722
FILING DATE:
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/456,647
FILING DATE: 02-JUN-1995
APPLICATION NUMBER: US 08/237,401
FILING DATE: 02-MAY-1994
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 07/884,486
FILING DATE: 15-MAY-1992
ATTORNEY/AGENT INFORMATION:
NAME: Wetherell Ph.D., John R.
REGISTRATION NUMBER: 31,678
```

```

REFERENCE/DOCKET NUMBER: 07251/007002
TELECOMMUNICATION INFORMATION:
    TELEPHONE: (619) 678-5070
    TELEFAX: (619) 678-5099
    INFORMATION FOR SEQ ID NO: 19:
        SEQUENCE CHARACTERISTICS:
            LENGTH: 3120 base pairs
            TYPE: nucleic acid
            STRANDEDNESS: single
            TOPOLOGY: linear
        MOLECULE TYPE: DNA
        IMMEDIATE SOURCE:
            CLONE: Tyro-10
        FEATURE:
            NAME/KEY: CDS
            LOCATION: 485..3047
US-09-158-722-19

Alignment Scores:
    Seq. No.:          39.4           Length:          3120
    Score:             56.00          Matches:         12
    Percent Similarity: 54.84%       Conservative:     5
    Best Local Similarity: 38.71%   Mismatches:      6
    Query Match:       29.02%       Indels:          8
    DB:                9           Gaps:            1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-158-722-19 (1-3120)
Oy      3 GlySerPheAlaLeuGlnAspSerPheSerSerLeuGlnGlyLeuGlyProGluTyr 22
Db      91 GGGAGCTGGCGCTCTTCACAGACTCA-----GCACCAGAGCAG 126
Oy      23 VallysLeuAluGlyLeuCysValCysLeuSer 33
Db      127 ATCTCATGTTTGGGCTGGATTGTGTCAAGC 159

RESULT 14
US-09-962-832-154
; Sequence 154: Application US//09962832
; Patent No. US20020110821A1
; GENERAL INFORMATION:
; APPLICANT: Ebner, Reinhard
; TITLE OF INVENTION: Cancer Gene Determination and Therapeutic Screening Using Signature
; FILE REFERENCE: 689290-74
; CURRENT APPLICATION NUMBER: US/09/962,832
; PRIORITY FILING DATE: 2001-09-25
; PRIOR APPLICATION NUMBER: US/60/235,077
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,280
; PRIOR FILING DATE: 2000-09-25
; NUMBER OF SEQ ID NOS: 259
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 154
; LENGTH: 302250
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-962-832-154

Alignment Scores:
    Pred. No.:          1.27e+04       Length:          302250
    Score:              56.00          Matches:         12
    Percent Similarity: 61.29%       Conservative:     7
    Best Local Similarity: 38.71%   Mismatches:      10
    Query Match:       29.02%       Indels:          2
    DB:                10           Gaps:            1

US-09-513-999C-7869_COPY_1_37 (1-37) x US-09-962-832-154 (1-302250)
Oy      7 LeuGlnAspSerPheSerSerLeuGlnGlyLeu-----LeuGlyProGluTyrVallys 24
Db      174135 CTTGAGGATCTGTGAATCCCTTAATTCGATGCAGACCTCTGGGGGTGTGGGTGTGTGTG 174194

```

[illegible]

TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL

COMMENT

Barrett, C., Burrows, J., Carter, M., Chacko, J., Chen, Z., Cox, C., David, R., Delgado, O., Deshazo, D., Ding, Y., Donah-Rashid, N., Dugan-Rocha, S., Durbin, K.J., Fernandez, C., Ferraguto, D., Forcman-Tansey, J., Frantz, P., Ganseth, R., Gorrell, J.H., Gorrell, L.L., Guervara, M., Harris, K., Hernandez, J., Hodgson, A., Hogues, M., Hollaway, C., Hosack, R., Jackson, L.E., Jackson, L., Jia, Y., Jones, M., Kelly, S., Kondejowski, N., Kong, Y., Kovar, C., Leal, B., Li, Z., Licharge, O., Liu, J., Liu, W., Logan, O., Lozano, R.J., Lu, J., Luchter, R., Mattin, R., Martinez, C., McLeod, M.P., Mel, G., Morgan, M., Morris, S., Nash, S., Nelson, A., Nguyen, R., Nguyen, N., Nguyen, S., Oswald, G., Parfitt, B., Patton, S., Patton, B., Perez, L., Pu, L., Quiles, R., Reiter, D., Rives, M., Sammel, S., Say, J., Scherer, S., Shah, E., Shen, H., Simon, M., Sparks, A., Stamps, A., Sugeng, S., Taber, P., Taylor, T., Vasquez, L., Vinson, R., Vo, O., Wahbah, M., Wallington, S., Weinstein, G., Weinstein, I.R., Williamson, A., Worley, K., Wren, J., Wrenford, G., Yu, W., Zhou, X., Nelson, D., and Gibbs, R.

Direct Submission
Unpublished
2 (bases 1 to 169620)
Worley, K.C.

Submitted (03-NOV-1999) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
On Aug 7, 2000 this sequence version replaced gi:8705345.

Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: <http://www.hgsc.bcm.tmc.edu/>
Contact: hgsc-help@bcm.tmc.edu

Project Information
Center project name: HMOG
Center clone name: RPI-458H3

Summary Statistics
Assembly program: Phrap; version 0.990329
Consensus quality: 139025 bases at least Q40
Consensus quality: 154842 bases at least Q30
Consensus quality: 159725 bases at least Q20
Estimated insert size: 162720; sum-of-contigs estimation
Estimated insert size: 171608; agarose-*rf* estimation
Quality coverage: 3.9x in Q20 bases; agarose-*rf* estimation
Quality coverage: 4.1x in Q20 bases; sum-of-contigs estimation

* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html)
* NOTE: This is a "working draft" sequence. It currently
* consists of 18 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

1	28689:	contig of 28689 bp in length
*	28690	gap of unknown length
*	28790	contig of 22043 bp in length
*	50833	gap of unknown length
*	50932:	gap of unknown length
*	50933	contig of 18212 bp in length
*	69145	gap of unknown length
*	69245	contig of 14960 bp in length
*	84205	gap of unknown length
*	84305	gap of 10365 bp in length
*	94668	gap of unknown length
*	94768	contig of 12494 bp in length
*	107261:	gap of unknown length
*	107361:	gap of unknown length
*	117550:	contig of 10189 bp in length
*	117551	gap of unknown length
*	117651	contig of 9289 bp in length
*	126939:	gap of unknown length
*	127040	contig of 8001 bp in length
*	135041	gap of unknown length
*	135141	contig of 6499 bp in length

BASE COUNT	ORIGIN	FEATURES
52024	a 33180 c 32128 g 50322 t 1966	others
		Location/Qualifiers
		1..169620
		/organism="Homo sapiens"
		/db_xref="taxon:9606"
		/chromosome="3"
		/clone="RP1-458H3"
		141640 141739: gap of unknown length
		141740 149558: contig of 7819 bp in length
		149559 149658: gap of unknown length
		149659 154562: contig of 4904 bp in length
		154563 154662: gap of unknown length
		154663 158987: contig of 4325 bp in length
		158988 159087: gap of unknown length
		159088 162376: contig of 3289 bp in length
		162377 162476: gap of unknown length
		162477 165191: contig of 2715 bp in length
		165192 165291: gap of unknown length
		165292 167173: contig of 1882 bp in length
		167174 167273: gap of unknown length
		167274 168393: contig of 1120 bp in length
		168394 168493: gap of unknown length
		168494 169620: contig of 1127 bp in length.

Query Match:	100.0%	Score 111:	DB 2:	Length 169620:
Best Local Similarity	100.0%	Pred.	0.6	5e-27:
Matches 111:	Conservative	0:	Mismatches	0:
			Indels	0:
			Gaps	0:

QY	1	ATGGGCGACATCTTTCGCTTCGAGGATCTTTTCATCTTTTGACGAGGACCTTCGGGGCCG	60
DB	87441	ATGGGCGACATCTTTCGCTTCGAGGATCTTTTCATCTTTTGACGAGGACCTTCGGGGCCG	87382
QY	61	GAGTATGTAACATCTCGGGCTCTGTGTGTGTGCTGAGTGGCTACTTACT	111
DB	87381	GAGTATGTAACATCTCGGGCTCTGTGTGTGTGCTGAGTGGCTACTTACT	87331

RESULT 2	
AC116565/C	
LOCUS	AC116565
DEFINITION	Homo sapiens chromosome 4 clone RP11-1263C18, WORKING DRAFT
VERSION	AC116565.3
KEYWORDS	HTG: HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEPIN.
SOURCE	Homo sapiens.

REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS
TITLE
JOURNAL
COMMENT

Eukariota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
1 (bases 1 to 199038)
Waterston, R.H.
The sequence of Homo sapiens clone
Unpublished
2 (bases 1 to 199038)
Waterston, R.H.
Direct Submission
Submitted (29-MAR-2002) Genome Sequencing Center, Washington
MO 63108, USA
4444 Forest Park Parkway, St. Louis,
MO 63108, USA
3 (bases 1 to 199038)
Waterston, R.H.
Direct Submission
Submitted (25-APR-2002) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
On Apr 25, 2002 this sequence version replaced gi:20128757.

----- Genome Center -----
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: <http://genome.wustl.edu/gsc/index.shtml>
Contact: submissions@wustl.wustl.edu

```

----- Project Information -----
Center project name: H_NH1263C18
----- Summary Statistics -----
Sequencing vector: M13, 0%
Sequencing vector: plasmid, 100%
Chemistry: Dye-Primer ET, 0% of reads
Chemistry: Dye-terminator Big Dye, 100% of reads
Assembly program: Phrap; version 0.990319
Consensus quality: 196905 bases at least Q40
Consensus quality: 197086 bases at least Q30
Consensus quality: 197234 bases at least Q20
Insert size: 20200; agarose-fp
Insert size: 19637; sum-of-ctrls
Quality coverage: 10.73 in Q20 bases; agarose-fp
Quality coverage: 10.82 in Q20 bases; sum-of-ctrls
-----
* NOTE: This is a 'working draft' sequence. It currently
* consists of 6 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 3283: contig of 3283 bp in length
* 3284 3383: gap of unknown length
* 3384 12884: contig of 9501 bp in length
* 12885 12984: gap of unknown length
* 12985 29821: contig of 16837 bp in length
* 29822 29922: gap of unknown length
* 29923 74050: contig of 44129 bp in length
* 74051 74151: gap of unknown length
* 74152 197985: contig of 123835 bp in length
* 197986 198086: gap of unknown length
* 198087 199038: contig of 953 bp in length.
* 199039
Location/Qualifiers
source
1. 199038
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="4"
/clone="RP11-1263C18"
misc_feature
1. 3283
/feature="assembly_name:Contig18"
3384. 12884
/feature="assembly_name:Contig19"
12985. 29821
/feature="assembly_name:Contig20"
29922. 74050
/feature="assembly_name:Contig21"
74151. 197985
/feature="assembly_name:Contig22"
198086. 199038
/feature="assembly_name:Contig14"
BASE COUNT 53552 a 46319 c 46377 g 52290 t 500 others
ORIGIN
Query Match 60.2% Score 66.8; DB 2; Length 199038;
Best Local Similarity 81.9%; Pred. No. 7,1e-12;
Matches 77; Conservative 0; Mismatches 17; Indels 0; Gaps 0;
QY 18 CTTGAGAGTTTCTTTTATCTTTGAGGAGGAGGAGTATGAAACTCCT 77
| | | | | | | | | | | | | | | | | | | | | | | | | |
Db 58755 CTGGAAGATGATTTCCACCTTCCAGGAGATCTCGGGCGGAGTATGTAACGCT 58696
| | | | | | | | | | | | | | | | | | | | | | | | | |
QY 78 GGGTCTGTGTGTCGTCGAGTGGCTGCTACT 111
| | | | | | | | | | | | | | | | | | | | | | | | | |
Db 58695 GGGTCTGTGTGTCGTCGAGTGGCTGCTGCT 58662
| | | | | | | | | | | | | | | | | | | | | | | | | |
RESULT 3
AC092586 AC092586 130636 bp DNA linear PRI 01-MAR-2002
LOCUS

```

```

DEFINITION Homo sapiens BAC clone RP11-30H15 from 2, complete sequence.
ACCESSION AC092586 AC016389
VERSION AC092586.2 GI:15638723
KEYWORDS
SOURCE
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 130636)
Sulston, J.E. and Waterston, R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
99063792
MEDLINE
9447074
REFERENCE
2 (bases 1 to 130636)
Haglund, K., Kozlowicz, A., Elliott, G. and Boyer, E.
The sequence of Homo sapiens BAC clone RP11-30H15
Unpublished (2001)
3 (bases 1 to 130636)
Waterston, R.H.
Direct Submission
Submitted (19-JUL-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
4 (bases 1 to 130636)
Waterston, R.H.
Direct Submission
Submitted (18-SEP-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
5 (bases 1 to 130636)
Waterston, R.
Direct Submission
Submitted (01-MAR-2002) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Sep 18, 2001 this sequence version replaced gi:14916171.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@watson.wustl.edu
----- Summary Statistics
Center project name: H_NH0030H15
Drafting Center: WIBR
-----

```

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:
The RP11-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Moon, P.Y., Zhao, B., Frengen, E., Teleno, M., Catanesi, J.J., and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at <http://www.chori.org>


```

KEYWORDS      HTG.
SOURCE        Homo sapiens.
ORGANISM      Homo sapiens
              Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
              Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE     1 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Genome Therapeutics Corporation Sequencing Center: Human Genome
              Sequence Data
JOURNAL       Unpublished
REFERENCE     2 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Direct Submission
JOURNAL       Submitted (25-JUN-2000) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
REFERENCE     3 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Direct Submission
JOURNAL       Submitted (14-JUL-2001) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
REFERENCE     4 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Direct Submission
JOURNAL       Submitted (28-MAR-2002) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
REFERENCE     5 (bases 1 to 140040)
AUTHORS       Smith,D.R.
TITLE         Direct Submission
JOURNAL       Submitted (09-APR-2002) Genome Therapeutics Corporation, 100 Beaver
              Street, Waltham, MA 02453, USA
COMMENT       On Mar 26, 2002 this sequence version replaced gi:14718295.

-----
Center: Genome Therapeutics Corporation
Center code: GTC
Web site: http://www.genomecorp.com/
Contact: gtc-seqcenter@genomecorp.com
-----
Project Information
Center project name: hg107

-----
IMPORTANT: This sequence is the entire
            Insert of clone RP11-93L14.
            The true right end of clone RP11-316G7 is at
            2000 in this sequence.

-----
Summary Statistics
Sequencing vector: N/A
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 990315
-----
Location/Qualifiers
1..140040
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="10"
/clone="RP11-93L14"
/clone_lib="RPCT-11"
BASE COUNT      48331 a 26787 c 24761 g 40161 t
ORIGIN
Query Match      58.2%; Score 64.6; DB 9; Length 140040;
Best Local Similarity 73.9%; Pred. No. 4e-11;
Matches 82; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
OY 1 ATGGGTGATCTTTTTCCTTCACAGATCTTTTCATCTTTGCACAGGACTCTTGAGGCG 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 94202 ATGTATGACCTCCCGCTTACATGATGTATGATACCTTTGGCCAGGATCCTGTGGCTG 94143
    Matches 82; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
OY 61 CAGTATGTAAACATCTCTGGTCTCTGTGTGTGTCCTGATGCTGCTTACT 111
    | ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 94142 GGGTATGTAAATCTCTGGTCTCTGTGTGTCATACCTGATGCTGCTGCT 94092

```

LOCUS	AL355812	147288 bp	DNA	linear	Pf1 13-JUN-2001							
DEFINITION	Human DNA sequence from clone RP11-81003 on chromosome Xq22.3-24, complete sequence.											
ACCESSION	AL355812											
VERSION	AL355812.23	GI:14456206										
KEYWORDS	HTG.											
SOURCE	human.											
ORGANISM	Homo sapiens											
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.											
AUTHORS	1 (bases 1 to 147288)											
TITLE	Smith M.											
JOURNAL	Direct Submission											
COMMENT	Submitted (13-JUN-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk requests: clonerequests@sanger.ac.uk On Jun 14, 2001 this sequence version replaced gi:14280423. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone configs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at http://www.sanger.ac.uk/HGP/chrX RP11-81003 is from the library RPC1-11.3 constructed by the group of Pletier de Jong. For further details see http://www.chori.org/bacpac/home.htm VECTOR: pBACe3.6 IMPORTANT: This sequence is not the entire insert of clone RP11-81003. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap. The true left end of clone RP11-81003 is at 1 in this sequence. The true left end of clone Rp6-204P4 is at 147189 in this sequence.											
FEATURES	Location/Qualifiers											
source	1. 147288											
	/organism="Homo sapiens"											
	/db_xref="taxon:9606"											
	/chromosome="X"											
	/map="q22.3-24"											
	/clone="RP11-81003"											
	/clone_lib="RPC1-11.3"											
repeat_region	1381..1674											
	/note="AluYb repeat: matches 1..296 of consensus"											
repeat_region	1757..1937											
	/note="AluYo repeat: matches 122..300 of consensus"											
repeat_region	2002..2645											
	/note="L1M4 repeat: matches 2110..2837 of consensus"											
repeat_region	2649..3323											
	/note="L1MC4 repeat: matches 6404..7210 of consensus"											
repeat_region	3811..3878											
	/note="L1MEC repeat: matches 2099..2180 of consensus"											
repeat_region	3882..4044											
	/note="L1ME3A repeat: matches 5665..5828 of consensus"											
repeat_region	4236..4533											
	/note="L1ME3A repeat: matches 5257..5563 of consensus"											
repeat_region	4724..5174											
	/note="L1M4 repeat: matches 4562..5026 of consensus"											
repeat_region	5401..5663											

repeat_region	/note="Charlie1a repeat: matches 1183. .1449 of consensus 5675. 5805
repeat_region	/note="L1PB3 repeat: matches 6014. .6145 of consensus" 5607. .6234
repeat_region	/note="l1 repeat: matches 4970. .5403 of consensus" 6251. .6998
repeat_region	/note="Charlie1a repeat: matches 404. .1196 of consensus" 6999. .7036
repeat_region	/note="19 copies 2 mer tt 100% conserved" 7244. .7864
repeat_region	/note="L1P repeat: matches 4160. .4781 of consensus" 7865. .8274
repeat_region	/note="Charlie1a repeat: matches 6. .417 of consensus" 8595. .8673
repeat_region	/note="MER58 repeat: matches 1. .83 of consensus" 8689. .8885
repeat_region	/note="L1M4 repeat: matches 5270. .5464 of consensus" 8942. .11399
repeat_region	/note="L1M4 repeat: matches 2661. .5149 of consensus" 11442. .11832
repeat_region	/note="L1M4 repeat: matches 2153. .2568 of consensus" 11837. .11994
repeat_region	/note="34 copies 2 mer tt 66% conserved" 11917. .12078
repeat_region	/note="L1ME2 repeat: matches 5989. .6155 of consensus" 12171. .13851
repeat_region	/note="L1MA9 repeat: matches 4533. .6275 of consensus" 13877. .14584
repeat_region	/note="L1M4 repeat: matches 5516. .6298 of consensus" 14582. .16120
repeat_region	/note="L1M1 repeat: matches 1441. .2965 of consensus" 16157. .16267
repeat_region	/note="L1M2 repeat: matches 1666. .1777 of consensus" 16278. .17065
repeat_region	/note="L1ME1 repeat: matches 5224. .6053 of consensus" 17066. .17368
repeat_region	/note="Alus repeat: matches 1. .302 of consensus" 17369. .17599
repeat_region	/note="L1ME1 repeat: matches 5010. .5224 of consensus" 17654. .19512
repeat_region	/note="L1MA1 repeat: matches 4389. .6262 of consensus" 19745. .20100
repeat_region	/note="L1MC2 repeat: matches 5470. .5826 of consensus" 20094. .20104
repeat_region	/note="L1 repeat: matches 4404. .4413 of consensus" 20105. .20666
repeat_region	/note="L1PA6 repeat: matches 5582. .6143 of consensus" 20667. .21348
repeat_region	/note="l1 repeat: matches 3734. .4404 of consensus" 21349. .21653
repeat_region	/note="Alus9 repeat: matches 1. .291 of consensus" 21634. .22811
repeat_region	/note="l1 repeat: matches 2511. .3734 of consensus" 22853. .23268
repeat_region	/note="L1M4 repeat: matches 3293. .3703 of consensus" 23269. .23420
repeat_region	/note="L1PA14 repeat: matches 6004. .6149 of consensus" 23422. .24030
repeat_region	/note="L1M4 repeat: matches 2655. .3293 of consensus" 24021. .24198
repeat_region	/note="L1M4 repeat: matches 2459. .2637 of consensus" 24207. .24312
repeat_region	/note="L1M4 repeat: matches -3. .103 of consensus" 24336. .24651
repeat_region	/note="L1MC6 repeat: matches 2291. .2280 of consensus" 24895. .25759
repeat_region	/note="L1MA9 repeat: matches 5440. .6304 of consensus" 25760. .26060
repeat_region	/note="Alus repeat: matches 1. .301 of consensus" 26061. .26155
repeat_region	/note="L1MA9 repeat: matches 5346. .5440 of consensus" 26213. .26559
repeat_region	/note="MIR1A1 repeat: matches 1. .354 of consensus" 26213. .26559

```

ACCESSION      AL591048
VERSION        AL591048.7   GI:15147695
KEYWORDS
SOURCE         human.
ORGANISM       Homo sapiens
REFERENCE      Enxaiyola; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS        Mammalia; Eutheria; Primates; Catarrhini; Homidae; Homo.
TITLE          1 (bases 1 to 75270)
JOURNAL
COMMENT        Direct Submission
Submitted (09-Aug-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
Requests: clonerequest@sanger.ac.uk
On Aug 10, 2001 this sequence version replaced gi:15020550.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence was finished as follows unless otherwise noted: all
regions were either double-stranded or sequenced with an alternate
chemistry or covered by high quality data (i.e., phred quality >=
30); an attempt was made to resolve all sequencing problems, such
as compressions and repeats; all regions were covered by at least
one plasmid subclone or more than one M13 subclone; and the
assembly was confirmed by restriction digest. The following
abbreviations are used to associate primary accession numbers given
in the feature table with their source databases: Em, EMBL; Sw,
SWISSPROT; Tr, TREMBL; Wp, WORMPEP. Information on the WORMPEP
database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
Chromosome 6, constructed by the Sanger Centre Chromosome 6 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/chr6
Rp11-612M16 is from the library RPCT-11.3 constructed by the group
of Pieter de Jong. For further details see
http://www.chori.org/dacpac/home.htm
VECTOR: pBACE3.6
IMPORTANT: This sequence is not the entire insert of clone
Rp11-612M16 It may be shorter because we sequence overlapping
sections only once, except for a 100 base overlap.
The true left end of clone Rp1-206F19 is at 73271 in this sequence.
The true right end of clone Rp11-82M9 is at 2000 in this sequence.
Location/Qualifiers
    source
        1..75270
            /organism="Homo sapiens"
            /db_xref="taxon:9606"
            /chromosome="6"
            /clone="RP11-612M16"
            /clone_lib="RPCT-11.3"
            /clone_1kb="RPCT-11.3"
            33923..34247
                /note="Sequence from reads from a short insert library
                derived from a single pUC clone. Restriction digest data
                confirm the assembly."
            46760..46849
                /note="Sequence from reads from a short insert library
                derived from a single pUC clone. Restriction digest data
                confirm the assembly."
            49353..49464
                /note="Sequence from reads from a short insert library
                derived from a single pUC clone. Restriction digest data
                confirm the assembly."
BASE COUNT     25749 a 14850 c 13893 g 20778 t
ORIGIN
Query Match    57.3%; Score 63.6; DB 9; Length 75270;
Best Local Similarity 73.6%; Pred. No. 8,7e-11;
Matches 81; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
1 ATGGGTGAGATCTTTGCCTTCAGAGATTCTTTTCATCTTGACAGGACTTCCTGGGCCG 60
||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

```

Df	1900		ATGTGCACAGCCTTCGCTTGGCTTGCCCTAAGCTGTGAACAATTTCCTGCAATCATCTCGGCGTG 1841
OY	61	GAGTATGTAAACCTCGGGTTCTGTGTGTGCCCTAGAOTGGCTGCTTAC 110 	
Df	1840	GAGTAGTAACACTTCTGGTCTGTGTCTGTCTGACCCTAGCACAGCTGTTCGC 1791	
RESULT 7			
Locus	AL137847/c		
DEFINITION	Homo sapiens	143372 bp	DNA linear PRI 16-NOV-2001
ACCESSION	AL137847		
VERSION	ALI37847.12	GI:16973786	
KEYWORDS	HTG.		
SOURCE	human. <i>Homo sapiens</i> Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.		
REFERENCE	AUTHORS TITLE JOURNAL		
COMMENT	<p>Direct Submission Submitted (16-NOV-2001) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humney@sanger.ac.uk Clone requests: clonerequests@sanger.ac.uk On Nov 17, 2001 this sequence version replaced g1:16408610. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: EM, EMBL, SW, SWISSPROT, Tr:, TREMBL, Mp:, WORMPEP. Information on the WORMPEP database can be found at http://www.sanger.ac.uk/projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping Group. Further information can be found at http://www.sanger.ac.uk/HGP/chrg Rp11-439k3 is from the library RPl1-11.2 constructed by the group of Pieter de Jong. For further details see http://www.chori.org/bacpac/home.htm VECTOR: pBACE3.6 IMPORTANT! This sequence is not the entire insert of clone Rpl1-439K3 It may be shorter because we sequence overlapping sections only once, except for a short overlap. The true left end of clone Rp11-439K3 is at 1 in this sequence. The true left end of clone Rp11-344I7 is at 14133 in this sequence.</p>		
FEATURES			
source	Location/Qualifiers 1..143372 /organism="Homo sapiens" /db_xref="taxon:9606" /chromosome="9" /map="q22.2-31.1" /clone="RP11-439K3" /clone_id="RPl1-11.2"		
BASE COUNT	42809 a	30224 c	28152 g 42187 t
ORIGIN			
Query Match	57.3%	Score 63.6:	DB 9: Length 143372;
Best Local Similarity	83.7%:	Pred.No. 8.7e-11;	
Matches	72: Conservative	0: Mismatched	14: Indels 0: Gaps 0:
Y	25	GATTCTTTTATCATCTTTCACGAGACTTCGTGGGCGCGAGTAGTATAAACCTCGTGGTCTC 84	

Db 82342 GATGAGATTTCCTCCCTGGCAGGAGATCCTGGGGCCAGAGTATGTAAACCTCTGGGCTTC 82283

QY 85 TGTGTGCTGCTGAGTGGCTGCTCTAC 110

Db 82282 TGTGTGCTGCTGAGTGGCTGCTCTGC 82257

RESULT 8	
LOCUS	AL137179/c
DEFINITION	AL137179 153053 bp DNA linear PRI 30-SEP-2000
ACCESSION	Human DNA sequence from clone RP11-82M9 on chromosome 6, complete sequence.
VERSION	AL137179
KEYWORDS	AL137179.14 GI:10443372
SOURCE	HTG.
ORGANISM	human.
	Homo sapiens
	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
	Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE	1 (bases 1 to 153053)
AUTHORS	Tracey/A.
TITLE	Direct Submission
JOURNAL	Submitted (26-SEP-2000) Sanger Centre, Hinxton, Cambridgeshire,

COMMENT

corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above. This sequence has been finished according to sequence map criteria as follows. An attempt is made to resolve all sequencing problems, such as compressions and repeats, but not necessarily within known annotated human repeat sequence elements (e.g. Alu). Where the sequence is ambiguous, there is an annotation using the 'unsure' feature key.

The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em1, EMBL; SW, SWISSPROT; Tr1, TREMBL; Wp1, WORMPEP; Information on the WORMPEP database can be found at <http://www.sanger.ac.uk/projects/C.elegans/wormpep> This sequence was generated from part of bacterial clone contigs of human chromosome 6, constituted by the Sanger Centre Chromosome 6 Mapping Group. Further information can be found at

RP11-82M9 is from the library RPCI-11 constructed at the Roswell Park Cancer Institute by the group of Pieter de Jong. For further details see <http://bacpac.med.buffalo.edu/VECTOR:pbac3.6>

IMPORTANT: This sequence is not the entire insert of clone RP11-82M9. It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap.

The true right end of clone RP11-82M9 is at 153053 in this sequence. The true right end of clone RP1-27919 is at 100 in this sequence.

FEATURES

Location/Qualifiers

```
source
1. 153053
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="6"
/clone="RP11-82w9"
/clone_lib="RPC1-11.1"
misc.feature
102234..102253
```

ORIGIN

Query Match	57.3%;	Score 63.6;	DB 9;	Length 153053;
Best Local Similarity	73.6%;	Pred. No. 8.7e-11;		

Matches	81: Conservative	0: Mismatches	29: Indels	0: Gaps	0:
Oy	1	ATGGGTGAGATCTTTTCTCTTGAGGATTCCTTTTATCTTTGACGGGACTTTGGGGCCG	60		
		ATGTCGACAGACTTTCCTCTGCTTCAGCGTAGACATATTTTGTCGTGATATCTCGGGGGCTG	152885		
Oy	61	GAGTATGTAAACCTCTGGGCTCTCTGTGTGTGTGCTCGAGTGGCTCTCTTAC	110		
		GAGTATGTAAAGCTTGTGGCTGTGTGTGTGTGCTCGACACACTCTTTTGC	152836		

RESULT	9
AC023408	
LOCUS	
DEFINITION	AC023408 153422 bp DNA linear HTG 07-JUN-2006
ACCESSION	Homo sapiens chromosome 6 clone RP11-612M16 map 6, WORKING DRAFT
VERSION	AC023408.22 unordered pieces.
KEYWORDS	AC023408.2 GI:8316895
SOURCE	HTG; HTGS_PHASE1; HTGS_DRAFT.
ORGANISM	Homo sapiens.
	Homo sapiens

REFERENCE	1 (bases 1 to 153422)
AUTHORS	Birtlen,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE	Homo sapiens chromosome 6, clone RP11-612M6
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 153422)
AUTHORS	Birtlen,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.

JOURNAL

COMMENT

Direct Submission
Submitted (14-FEB-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Jun 7, 2000 this sequence version replaced gi:6970573.
All repeats were identified using RepeatMasker:
Smith, A.F.A. & Green, P. (1996-1997)

----- Genome Center
http://ftp.genome.washington.edu/RM/RepeatMasker.html

Center: Whitehead Institute/ MIT Center for Genome Research

Center code: WIBK

web site: <http://www-seq.wi.mit.edu>

contact: sequence_submissions@ncbi.nlm.nih.gov

***** Project Information *****
Contract number: 16363

Center project name: E0303
Center phone name: 612 M 16

----- Summary Statistics -----

Sequencing vector: M13: 100% of reads

Chemistry: Dye-terminator Big Dye: 100% of reads

Assembly program: Phrap: version 0.960731

Consensus quality: 135238 bases at least 040

Consensus quality: 141969 bases at least Q30

Consensus quality: 146244 bases at least Q20

Insert size: 150000; agarose-fp

Insert size: 151322; sum-of-contigs

Quality coverage: 4.0 in Q20 bases; agarose-1p
Quality coverage: 3.9 in Q20 bases; sum-of-continfs

NOTE: This is a 'working draft' sequence. It currently consists of 22 continfs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the continfs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

```
1 1152: continf of 1152 bp in length
* 1153 1252: gap of 100 bp
* 1253 2252: continf of 1000 bp in length
* 2253 2352: gap of 100 bp
* 2353 3503: continf of 1151 bp in length
* 3504 3603: gap of 100 bp
* 3604 4658: continf of 1055 bp in length
* 4659 4758: gap of 100 bp
* 4759 6059: continf of 1301 bp in length
* 6060 6159: gap of 100 bp
* 6160 6832: continf of 673 bp in length
* 6833 6932: gap of 100 bp
* 6933 9547: continf of 2615 bp in length
* 9548 9647: gap of 100 bp
* 9648 11267: continf of 1620 bp in length
* 11268 11367: gap of 100 bp
* 11368 14730: continf of 3363 bp in length
* 14731 14830: gap of 100 bp
* 14831 18501: continf of 3671 bp in length
* 18502 18601: gap of 100 bp
* 18602 23876: continf of 5275 bp in length
* 23877 23976: gap of 100 bp
* 23977 28125: continf of 4149 bp in length
* 28126 28225: gap of 100 bp
* 28226 33384: continf of 5159 bp in length
* 33385 33484: gap of 100 bp
* 33485 37950: continf of 4466 bp in length
* 37951 38050: gap of 100 bp
* 38051 43278: continf of 5228 bp in length
* 43279 43378: gap of 100 bp
* 43379 49641: continf of 6263 bp in length
* 49642 49741: gap of 100 bp
* 49742 57239: continf of 7498 bp in length
* 57240 57339: gap of 100 bp
* 57340 65126: continf of 7787 bp in length
* 65127 65226: gap of 100 bp
* 65227 74642: continf of 9416 bp in length
* 74643 74742: gap of 100 bp
* 74743 86960: continf of 12218 bp in length
* 86961 87060: gap of 100 bp
* 87061 115886: continf of 28826 bp in length
* 115887 115987: gap of 100 bp
* 115987 153422: continf of 37436 bp in length.
Location/Qualifiers
source
1. 153422
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="6"
/map="6"
/clone="RP11-612M16"
/clone_lib="RP11 Human Male BAC"
1. 1152
/misc_feature
/note="assembly_fragment"
1253. 2252
/misc_feature
/note="assembly_fragment"
2353. 3503
/misc_feature
/note="assembly_fragment"
3604. 4658
/misc_feature
/note="assembly_fragment"
4759. 6059
/misc_feature
/note="assembly_fragment"
6160. 6832
```

```
/note="assembly_fragment
clone_end:17
vector_side:right"
misc_feature
6933. 9547
/note="assembly_fragment"
9648. 11267
misc_feature
11368. 14730
/note="assembly_fragment"
14831. 18501
/note="assembly_fragment"
18602. 23876
/misc_feature
23977. 28125
/note="assembly_fragment"
28226. 33384
/misc_feature
33485. 37950
/note="assembly_fragment"
38051. 43278
/note="assembly_fragment
clone_end:SP6
vector_side:right"
43379. 49641
/note="assembly_fragment"
49742. 57239
/note="assembly_fragment"
57340. 65126
/note="assembly_fragment"
65227. 74642
/note="assembly_fragment"
74743. 86960
/misc_feature
87061. 115886
/note="assembly_fragment"
115987. 153422
/note="assembly_fragment"
misc_feature
BASE COUNT 47446 a 29388 c 28599 g 45687 t 2102 others
ORIGIN
Query Match 57.3%; Score 63.6; DB 2; Length 153422;
Best Local Similarity 73.6%; Pred. No. 8.7e-11;
Matches 81; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
QY 1 ATGGCTGATCTTTGCTTCAGAGATCTTTTCATCTTCAGAGACTTGCGGCG 60
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 49270 ATGCGAGACCTCTTGCTTCAGAGACTTCATCTTCAGAGACTTGCGGCG 49329
QY 61 GAGTATGTAACCTCGGCTCTGTCGCTGAGTGCCTGCTCTAC 110
||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 49330 GAGTATGTAACCTCGGCTCTGTCGCTGAGTGCCTGCTCTAC 49379
RESULT 10
AC109592/c 86314 bp DNA linear PRI 13-MAR-2002
LOCUS Homo sapiens BAC clone RP11-750022 from 4, complete sequence.
DEFINITION AC109592
ACCESSION AC109592.3 GI:19310357
VERSION
KEYWORDS HTG.
SOURCE Homo sapiens.
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
REFERENCE 1 (bases 1 to 86314)
Sulston, J.E. and Waterston, R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
JOURNAL MEDLINE
PUBMED 99063792
9847074
2 (bases 1 to 86314)
REFERENCE Buatsi, D. and Kozlowicz, A.
The sequence of Homo sapiens BAC clone RP11-750022
AUTHORS
TITLE
```

JOURNAL Unpublished (2001)
REFERENCE 3 (bases 1 to 86314)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (05-FEB-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE 4 (bases 1 to 86314)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (06-MAR-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE 5 (bases 1 to 86314)
AUTHORS Waterston,R.H.
TITLE Direct Submission
JOURNAL Submitted (09-MAR-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE 6 (bases 1 to 86314)
AUTHORS Waterston,R.
TITLE Direct Submission
JOURNAL Submitted (13-MAR-2002) Department of Genetics, Washington University 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA On Mar 9, 2002 this sequence version replaced gi:18642965.

----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: saplens@watson.wustl.edu

----- Summary Statistics
Center project name: H_NH0750022

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted:
all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:
Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see http://genome.wustl.edu/gsc

SOURCE INFORMATION:
The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osoegawa,K., Moon,P.Y., Zhao,B., Frengen,E., Tateno,M., Catanese,J.J. and de Jong,P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (http://www.resgen.com) or Pieter de Jong and coworkers at http://www.chori.org
VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:
The clone sequenced to the left is RP11-510D4, 2000 bp overlap; the clone sequenced to the right is RP11-36611, 2000 bp overlap. Actual start of this clone is at base position 149897 of RP11-510D4; actual end is at base position 44359 of RP11-36611.

FEATURES

Source
1. 86314
/organism="Homo sapiens"
/db_xref="taxon:9606"
/chromosome="4"

```

/map="4"
/clone="RP11-750022"
/clone_id="RPCI-11"
2..160
/rpt_family="Alu"
repeat_region
256..655
/rpt_family="L1"
repeat_region
954..1182
/rpt_family="L1"
repeat_region
1216..1472
/rpt_family="L1"
repeat_region
1563..1876
/rpt_family="L1"
repeat_region
2896..3255
/rpt_family="MaLR"
repeat_region
3256..4669
/rpt_family="MaLR"
repeat_region
4670..4969
/rpt_family="MaLR"
repeat_region
10088..10184
/rpt_family="MaLR"
repeat_region
11452..11617
/rpt_family="CRL"
repeat_region
12716..12930
/rpt_family="CRL"
repeat_region
13258..13395
/rpt_family="MIR"
repeat_region
14378..14415
/rpt_family="AT-rich"
repeat_region
14817..14883
/rpt_family="Alu"
repeat_region
15540..15842
/misc_feature
note="match to EST AI023549 (NID:g3238593) ov79f12.s1"
21639..21945
/rpt_family="L1"
repeat_region
23095..23142
/rpt_family="(TG)n"
repeat_region
24837..24987
/rpt_family="ERV1"
repeat_region
24994..26519
/rpt_family="ERV1"
repeat_region
26550..26833
/rpt_family="ERV1"
repeat_region
26834..26918
/rpt_family="(TA)n"
repeat_region
26938..26973
/rpt_family="(GA)n"
repeat_region
26979..27252
/rpt_family="Alu"
repeat_region
27546..27618
/rpt_family="MaLR"
repeat_region
27984..28617
/rpt_family="L1"
repeat_region
28624..28942
/rpt_family="Alu"
repeat_region
29073..29150
/rpt_family="MER53"
repeat_region
29081..29158
/rpt_family="MER53"
repeat_region
29199..29562
/rpt_family="L1"
repeat_region
29563..30039
/rpt_family="L1"
repeat_region
30313..31555
/rpt_family="L1"
repeat_region
31554..31738
/rpt_family="L1"
repeat_region
31739..31786
/rpt_family="AT-rich"
repeat_region
34024..34352
/rpt_family="Alu"
repeat_region
38180..38776
/rpt_family="L1"

```

Query Match	57.1%;	Score 63.4;	DB 9;	Length 86314;
Best Local Similarity	82.0%;	Pred. No. 1e-10;		
Matches 73; Conservative	0;	Mismatches 16;	Indels 0;	Gaps 0;

32222 TCTGTGTGCCCCAGTGCGTCTGCT 32194

RESULT	11
AC021378/c	
LOCUS	
DEFINITION	Homo sapiens clone RP11-26G6, WORKING DRAFT SEQUENCE, 10 unordered pieces.
ACCESSION	AC021378
VERSION	AC021378.4 GI:8072570
KEYWORDS	HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE	Homo sapiens.
ORGANISM	Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE	1 (bases 1 to 159475) Birren,B., Linton,L., Nusbaum,C. and Lander,E. Homo saplens, clone RPl1-26G6 Unpublished 2 (bases 1 to 159475)
AUTHORS	Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N., Anderson,S., Baldwin,J., Barna,N., Beckerly,R., Bega,F., Boudslavsky,I., Boukhgalter,B., Brown,A., Bukett,G., Castle,A., Chopel,Y., Colangelo,M., Collins,S., Collimore,A., Cooke,P., Deatellano,K., Dewar,K., Domino,M., Doyle,W., Fenesfor,J., Ferrelita,P., Fitzhugh,W., Forrest,C., Gage,D., Galagan,J.,
JOURNAL	
REFERENCE	
AUTHORS	

COMMENT

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: <http://www.seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
Project Information

----- Summary Statistics

Assembly program: Phrap; version 0.960731

consensus quality: 150152 bases at least
consensus quality: 155379 bases at least

Consensus quality: 157415 bases at least Q20
 Consensus quantity: 157400 bases at least Q30

Insert size: 157000; agarose-gel
Insert size: 158575; sum-of-contigs

Quality coverage: 4.2 in Q20 bases; agarose-fp
Quality coverage: 4.2 in Q20 bases; sum-of-contigs

NOTE: This is a 'working draft' sequence. It currently consists of 10 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of 'N', but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

*	1	2335:	contlg of 2315 bp in length
*	216	2445:	gap of 100 bp
*	2416	9041:	contlg of 6626 bp in length
*	9042	9141:	gap of 100 bp
*	9142	147359:	contlg of 8218 bp in length
*	17360	17459:	gap of 100 bp
*	17460	246087:	contlg of 8608 bp in length
*	26068	26187:	gap of 100 bp
*	26168	40876:	contlg of 14709 bp in length
*	40877	40996:	gap of 100 bp
*	40927	556845:	contlg of 15869 bp in length
*	55646	556945:	gap of 100 bp
*	74625	74524:	contlg of 17579 bp in length
*	74625	74624:	gap of 100 bp
*	94603	94662:	contlg of 20038 bp in length
*	94603	94762:	gap of 100 bp
*	94763	1269545:	contlg of 32163 bp in length
*	126926	127025:	gap of 100 bp
*	127026	159475:	contlg of 32450 bp in length

Source

misc_feature

```

/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="RP11-2666"
/clone_lib="RPCI-11 Human Male BAC
1.2315
/note="assembly_fragment

```

```

misc_feature      clone.end:SP6
                  vector.side:left"
misc_feature      2416..9041
                  /note="assembly_fragment"
misc_feature      9142..17359
                  /note="assembly_fragment"
misc_feature      17460..26067
                  /note="assembly_fragment"
misc_feature      26168..40876
                  /note="assembly_fragment"
misc_feature      40977..56845
                  /note="assembly_fragment"
misc_feature      56946..74524
                  /note="assembly_fragment"
misc_feature      74625..94662
                  /note="assembly_fragment"
misc_feature      94763..126925
                  /note="assembly_fragment"
misc_feature      127026..159475
                  /note="assembly_fragment"
misc_feature      /note="assembly_fragment"
                  clone.end:T7
                  vector.side:left"
BASE COUNT      52037 a 27831 c 28355 g 50348 t 904 others
ORIGIN

```

```

Query Match      57.1% Score 63.4: DB 2: Length 159475;
Best Local Similarity 82.0% Pred. No. 1e-10;
Matches 73: Conservative 0; Mismatches 16; Indels 0; Gaps 0;

```

```

Db 23 AGAATCTTTTTCATCTTTCGAGGACTTCTGCGCGGAGTATGTAACCTCTGGGTC 82
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Db 45765 AGGATGATCTTCTGCTGCGGGAATCTGCGGCGTATGTAACCTCTGGGTC 45706
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Qy 83 TCTGTGTGCTGCTAGTGTGCTGCTACT 111
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||
Db 45705 TCTGTGTGCTGCTGCTGCTGCTGCT 45677
    |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||  |||||

```

```

RESULT 12
LOCUS      AL157937 153783 bp DNA linear PRI 04-APR-2001
DEFINITION Human DNA sequence from clone RP11-408N14 on chromosome 9 Contains
ACCESSION AL157937
VERSION   AL157937.21 GI:11121006
KEYWORDS  HMC.
SOURCE    human.
ORGANISM  Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
REFERENCE 1 (bases 1 to 153783)
AUTHORS   Kimberley A.
JOURNAL   Direct Submission
Submitted (26-FEB-2001) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquerry@sanger.ac.uk
Requests: clonerequests@sanger.ac.uk
On Nov 8, 2000 this sequence version replaced gi:11071307.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
validation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
The following abbreviations are used to associate primary accessions
numbers given in the feature table with their source databases:
Em, EMBL; Sw, SWISSPROT; Tr, TREMBL; Wp, WORMPEP; Information
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone configs of human
chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr9
IMPORTANT: This sequence is not the entire insert of clone

```

RP11-408N14 It may be shorter because we sequence overlapping sections only once, except for a 100 base overlap. The true left end of clone RP11-408N14 is at 1 in this sequence. The true left end of clone RP11-44115 is at 13684 in this sequence. The true right end of clone RP11-145E5 is at 29400 in this sequence. This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. RP11-408N14 is from the library RP11-11.2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>

FEATURES

source

```

repeat_region    /note="MIR repeat: matches 21..190 of consensus"
repeat_region    499..941
                  /note="MIR repeat: matches 4921..5372 of consensus"
repeat_region    1151..1457
                  /note="AluMb repeat: matches 1..304 of consensus"
repeat_region    1671..2296
                  /note="L1MC5 repeat: matches 7249..7897 of consensus"
repeat_region    2401..2642
                  /note="L1MC5 repeat: matches 6983..7227 of consensus"
repeat_region    3314..3359
                  /note="23 copies 2 mer tg 93% conserved"
repeat_region    3769..3991
                  /note="MIR repeat: matches 5..241 of consensus"
repeat_region    5093..5394
                  /note="match: GSS: Em:AQ823856"
repeat_region    5306..5391
                  /note="match: GSS: Em:AQ823856"
repeat_region    5537..5881
                  /note="L2 repeat: matches 2164..2262 of consensus"
repeat_region    5607..5973
                  /note="MLT2FA repeat: matches 5..324 of consensus"
repeat_region    6531..7177
                  /note="match: GSS: Em:AQ137344"
repeat_region    6538..6935
                  /note="match: GSS: Em:B91342"
repeat_region    6611..6910
                  /note="match: GSS: Em:AQ217550"
repeat_region    7761..7811
                  /note="AluIo repeat: matches 1..297 of consensus"
repeat_region    7812..8120
                  /note="L2 repeat: matches 2651..2694 of consensus"
repeat_region    8121..8174
                  /note="AluXk repeat: matches 1..309 of consensus"
repeat_region    8950..9125
                  /note="L2 repeat: matches 2576..2651 of consensus"
repeat_region    9360..9878
                  /note="match: GSS: Em:AQ44387"
repeat_region    9753..9824
                  /note="match: GSS: Em:AQ771822"
repeat_region    9869..9980
                  /note="36 copies 2 mer tg 79% conserved"
repeat_region    11255..17292
                  /note="MIR repeat: matches 139..262 of consensus"
repeat_region    17910..18096
                  /note="L1Pa5 repeat: matches 76..6143 of consensus"
repeat_region    /note="L1MC4 repeat: matches 7641..7835 of consensus"

```


repeat_region	16520..20148	/note="L1P12 repeat: matches 1421..1009 of consensus"
repeat_region	20127..21370	/note="L1P11 repeat: matches 921..2161 of consensus"
repeat_region	21372..21581	/note="L1P11 repeat: matches 2262..6165 of consensus"
repeat_region	25293..25713	/note="L1P10 repeat: matches 5655..6083 of consensus"
repeat_region	26120..26412	/note="L1P12 repeat: matches 1541..1650 of consensus"
misc_feature	26198..26577	/note="match: GSS: Em:AQ147782"
repeat_region	26660..26962	/note="L1P12 repeat: matches 1..300 of consensus"
misc_feature	complement(27055..27448)	/note="match: STS: Em:HSJ37G11"
repeat_region	27407..31040	/note="L1P12 repeat: matches 2505..6164 of consensus"
repeat_region	31031..31138	/note="L1P repeat: matches 1541..1650 of consensus"
repeat_region	31144..31497	/note="HEIB repeat: matches 1..364 of consensus"
repeat_region	31498..33130	/note="HEIB-INTERNAL repeat: matches 1..1580 of consensus"
repeat_region	33131..33479	/note="HEIB repeat: matches 1..359 of consensus"
repeat_region	33485..35130	/note="L1P13 repeat: matches 248..1557 of consensus"
repeat_region	36624..36669	/note="23 copies 2 mer ta 76% conserved"
repeat_region	36680..36897	/note="L1MCA repeat: matches 6404..6623 of consensus"
repeat_region	37335..37508	/note="37 copies 2 mer ag 75% conserved"
repeat_region	38923..39151	/note="Charlie4 repeat: matches 231..1926 of consensus"
misc_feature	complement(39281..39766)	/note="match: GSS: Em:AQ894025"
repeat_region	40317..40540	/note="MR repeat: matches 20..234 of consensus"
repeat_region	40603..40726	/note="L2 repeat: matches 2587..2710 of consensus"
repeat_region	43871..43952	/note="L2 repeat: matches 2673..2750 of consensus"
repeat_region	43962..44259	/note="AluY repeat: matches 1..299 of consensus"
repeat_region	45076..45280	/note="MR repeat: matches 2..205 of consensus"
repeat_region	45413..45588	/note="MR repeat: matches 1..205 of consensus"
misc_feature	complement(46149..46586)	/note="match: GSS: Em:AQ343245"
repeat_region	47817..47938	/note="L2 repeat: matches 2627..2749 of consensus"
repeat_region	48349..48579	/note="MR11 repeat: matches 1..220 of consensus"
repeat_region	50262..50664	/note="L2 repeat: matches 2339..2749 of consensus"
repeat_region	50674..52080	/note="L1M1 repeat: matches 1389..30 of consensus"
repeat_region	52084..52853	/note="L1M4 repeat: matches 5516..6308 of consensus"
repeat_region	54498..60638	/note="L1P4 repeat: matches 1..6146 of consensus"
repeat_region	61367..61560	/note="MR1P repeat: matches 346..541 of consensus"
repeat_region	62305..62430	/note="63 copies 2 mer tt 59% conserved"
repeat_region	63089..63629	/note="L1P12 repeat: matches 10..540 of consensus"
repeat_region	63630..63887	/note="AluY repeat: matches 7..263 of consensus"

repeat_region	63888..64059	/note="A1US3 repeat: matches 134..303 of consensus"
repeat_region	64060..64352	/note="A1US3 repeat: matches 1..295 of consensus"
repeat_region	64353..64485	/note="A1US3 repeat: matches 1..134 of consensus"
repeat_region	64486..64834	/note="A1US3 repeat: matches 539..-291 of consensus"
repeat_region	64835..64938	/note="A1PA10 repeat: matches 1761..1897 of consensus"
repeat_region	64939..65350	/note="A1R14B repeat: matches 1..608 of consensus"
repeat_region	65351..67266	/note="A1PA10 repeat: matches 1894..3818 of consensus"
repeat_region	67317..69162	/note="A1PA10 repeat: matches 3793..5643 of consensus"
repeat_region	69122..69167	/note="A1PA10 repeat: matches 5616..5662 of consensus"
repeat_region	69216..69754	/note="A1PA10 repeat: matches 5610..6165 of consensus"
repeat_region	70283..70389	

Query Match:	56.8%	Score 63:	DB 9:	Length 153783:
Best Local Similarity:	73.0%	Pred. No. 1,4e-10:		
Matches 81:	Conservative	0:	Mismatches 30:	Indels 0:
			Gaps	0:
QY	1	ATGGGTCGATCTTTGCCCTGCAGATCTTTTATCTTTTGCAGGAGCACTTCGGGGCCG	60	
Db	18987	ATGACAGACATCTCCCTCTGCTCTAGTATGCAGAACATCTTCTGGGGATCTCGGGGCGG	18928	
QY	61	GAGTATGTAACCTCCGGGCTCTGTGTGTGTCCTGAGTGGCTGCTCTACT	111	
Db	18927	GAGTATGTAAGCTCCGCGGCTCTGTGTGTGTCATCAATCCAGACACACTCTCTGCT	18877	

RESULT 13	AC021025/c	standard; DNA; HTG: 123779 BP.
ID	AC021025	
XX		
AC	AC021025;	
XX		
SV	AC021025.9	
XX		
DT	14-JAN-2000 (Rel. 62, Created)	
DT	08-AUG-2000 (Rel. 64, Last updated, Version 11)	
XX		
DE	Homo sapiens chromosome 3 clone RP11-79K17, WORKING DRAFT SEQUENCE, 10	
DE	unordered pieces.	
XX		
KW	HTG: HTGS_DRAFT; HTGS_PHASE1.	
XX		
OS	Homo sapiens (human)	
OC	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia;	
OC	Eutheria; Primates; Catarrhini; Hominiidae; Homo.	
XX		
RM	[1]	
RP	1-123779	
RA	Muzny D.M., Adams C., Bailey M., Barbara J., Blankenburg K., Bodota B.,	
RA	Bouck Y., Bowle S., Brooks A., Buhay C., Bunac C., Burkett C., Burrows J.,	
RA	Catter M., Chacko J., Chen Z., Cox C., David R., Delgado O., Deshazo D.,	
RA	Ding Y., Domah-Nashid N., Dugan-Rocha S., Dublin K.J., Fernandez C.,	
RA	Ferrazuto D., Forcum-Thaneey J., Frintz P., Ganesh R., Gorrell J.H.,	
RA	Gorrell L.L., Guevara W., Harris K., Hernandez J., Hodgson A., Hogues M.,	
RA	Holloway C., Hosak H., Jackson L.E., Jackson L., Jia Y., Jones M.,	
RA	Kelly S., Kondelewski N., Kong Y., Kovar C., Leal B., Li Z., Lichtarge O.,	
RA	Lin J., Liu W., Logan O., Lozada R.J., Lu J., Lucier R., Martin R.,	
RA	Matlinetz C., McLeod M.P., Mei G., Morgan M., Morris S., Nash S., Nelson A.,	
RA	Nguyen R., Nguyen N., Nguyen S., Oswal G., Parish B., Paxton S., Payton B.,	
RA	Perez L., Pu L.L., Quiles M., Reiter D., Rives M., Samuel S., Say J.,	
RA	Scherer S., Shah E., Shen H., Simon M., Sparks A., Stamps A., Sugang R.,	
RA	Tabor P., Taylor T., Vasquez L., Vinson R., Vo O., Wahbah M.,	
RA	Wellington S., Weinstock G., Weinstock I.R., Williamson A., Worley K.,	
RA	Wren J., Wrenford G., Yu W., Zhou X., Nelson D., Gibbs R.:	

GenCore version 5.1.4-p5_4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 00:52:59 ; Search time 150 seconds

(without alignments)
1666.479 Million cell updates/sec

Title: US-09-513-999C-3792_COPY_51_161

Perfect score: 1 atgggtgacatttcctt.....gcctgagtgcgtctact 111

Sequence: IDENTITY NUC

Gapop 10.0, Gapext 1.0

Scoring table: 2185239 segs, 112599159 residues

Searched: 1 number of hits satisfying chosen parameters: 4370478

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : _N_Geneseq_101002.*

1: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1980.DAT:*
2: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1981.DAT:*
3: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1982.DAT:*
4: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1983.DAT:*
5: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1984.DAT:*
6: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1985.DAT:*
7: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1986.DAT:*
8: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1987.DAT:*
9: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1988.DAT:*
10: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1989.DAT:*
11: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1990.DAT:*
12: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1991.DAT:*
13: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1992.DAT:*
14: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1993.DAT:*
15: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1994.DAT:*
16: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1995.DAT:*
17: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1996.DAT:*
18: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1997.DAT:*
19: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1998.DAT:*
20: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA1999.DAT:*
21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT:*
22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT:*
23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT:*
24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	447	21	AAC03794 Human secreted pro
2	111	100.0	447	21	AAZ42680 Human 5' EST isola
3	58.2	52.4	349	22	AAE5888 Novel human polynu
4	58.2	52.4	570	22	ABA63453 Human foetal liver
5	58.2	52.4	570	22	ABA30652 Probe #9118 for ge
6	58.2	52.4	570	22	AAK11985 Human brain expres
7	58.2	52.4	570	22	AAK37688 Human bone marrow
8	58.2	52.4	570	22	AA118447 Probe #8380 for ge
9	58.2	52.4	570	22	AA135563 Probe #12249 used

C 10	58.2	52.4	570	24	ABS11680 Human genome-deriv
C 11	56.4	50.8	128600	24	ABK83461 Human cDNA differe
C 12	55	49.5	1982	21	AAK68089 Human secreted pro
C 13	52.8	47.6	609	22	AAH98980 Human EST-derived
C 14	50.8	45.8	1299	22	AAH91827 DNA encoding novel
C 15	46	41.4	807	23	AAH64647 DNA encoding novel
C 16	46	41.4	843	23	AAH79254 DNA encoding novel
C 17	44	39.6	1996	22	ABA18896 Human nervous syst
C 18	43.8	39.5	1909	23	AAH88434 DNA encoding novel
C 19	40.2	36.2	240	24	AAH25478 Human ORX polynuc
C 20	39	35.1	660	23	AAH73441 DNA encoding novel
C 21	38.8	35.0	162	22	ABA75762 Human foetal liver
C 22	38.8	35.0	519	22	ABA63320 DNA encoding novel
C 23	36.2	32.6	2787	23	AAH68019 DNA encoding novel
C 24	36.2	32.6	2787	23	AAH68019 DNA encoding novel
C 25	35.2	31.7	1173	23	AAH69200 DNA encoding novel
C 26	35.2	31.7	1173	23	AAH82303 DNA encoding novel
C 27	35.2	31.7	1173	23	AAH86064 DNA encoding novel
C 28	35.2	31.7	1291	23	AAH77610 DNA encoding novel
C 29	35.2	31.7	2097	23	AAH77443 DNA encoding novel
C 30	35.2	31.7	2569	23	AAH77685 DNA encoding novel
C 31	35.2	31.7	11534	24	ABL32342 Human immune syste
C 32	34.2	30.8	1278	23	AAH72474 DNA encoding novel
C 33	33.4	30.1	396	22	AAH74465 Human immune/haema
C 34	32.4	29.2	447	22	AAH16101 Human breast cance
C 35	32.4	29.2	447	22	AAH124945 Human breast cance
C 36	31.8	28.6	349980	22	AAH86431 Pyrococcus abyssi
C 37	31.8	28.6	349980	22	AAH41223 Pyrococcus abyssi
C 38	31	27.9	826	22	AAH04786 Human cDNA clone (
C 39	30.8	27.7	3659	7	AAH60204 Interleukin-pseudo-
C 40	30.6	27.6	352	22	AAH59083 Human immune/haema
C 41	29.2	26.3	369	24	AAH64361 Human benign prost
C 42	29.2	26.3	369	24	ABL61736 Colon adenocarcino
C 43	29.2	26.3	369	24	ABL65612 Lung cancer relate
C 44	29.2	26.3	369	24	ABL66228 Lung cancer relate
C 45	29.2	26.3	1314	22	AAH31530 Human DNA for a no

ALIGNMENTS

RESULT 1	
AAC03794	AAC03794 standard; cDNA; 447 BP.
XX	XX
XX	XX
AC	AAC03794:
XX	XX
DT	06-OCT-2000 (first entry)
XX	XX
DE	Human secreted protein 5' EST, SEQ ID NO: 3792.
XX	XX
KW	Human: 5' EST: expressed sequence tag; secreted protein; cDNA isolation;
KW	gene therapy; chromosome mapping; ss.
XX	XX
OS	Homo sapiens.
XX	XX
PN	EP1033401-A2.
XX	XX
PD	06-SEP-2000.
XX	XX
PF	21-FEB-2000; 2000EP-0200610.
XX	XX
PR	26-FEB-1999; 99US-0122487.
XX	XX
PA	(GEST) GENSET.
XX	XX
PI	Dumas Milne Edwards J, Duclert A, Giordano J;
XX	XX
DR	WPI; 2000-500381/45.
XX	XX
PT	P-PADB; AAC03788.
XX	XX
PT	New nucleic acid that is a 5' expressed sequence tag (5' EST) for
XX	obtaining cDNAs and genomic DNAs that correspond to 5'ESTs and for

PT diagnostic, forensic, gene therapy and chromosome mapping procedures -
 XX
 PS Claim 1: SEQ ID 3792; 71pp + CD-ROM; English.
 XX
 CC The present sequence is one of a large number of 5' ESTs derived from
 CC mRNAs encoding secreted proteins. An ORF has been identified within the
 CC sequence. The 5' ESTs were prepared from total human RNAs or polyA+ RNAs
 CC derived from 30 different tissues. EST sequences usually correspond
 CC mainly to the 3' untranslated region (UTR) of the mRNA because they are
 CC often obtained from oligo-dT primed cDNA libraries. Such ESTs are not
 CC well suited for isolating cDNA sequences derived from the 5' ends of
 CC mRNAs and even in those cases where longer cDNA sequences have been
 CC obtained, the full 5' UTR is rarely included. 5' ESTs are derived from
 CC mRNAs with intact 5' ends and can therefore be used to obtain full length
 CC cDNAs and genomic DNAs. 5' ESTs are also used in diagnostic, forensic,
 CC gene therapy and chromosome mapping procedures. They are used to obtain
 CC upstream regulatory sequences and to design expression and secretion
 CC vectors.
 CC
 CC Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;
 CC
 CC Query Match 100.0%; Score 111; DB 21; Length 447;
 CC Best Local Similarity 100.0%; Pred. No. 1.4e-27;
 CC Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 CC
 CC QY 1 ATGGGTGATCTTTGCGTGCAGATTCCTTTTCATCTTTGACAGGACTTCGGGCGC 60
 CC |
 CC DB 51 ATGGGTGATCTTTGCGTGCAGATTCCTTTTCATCTTTGACAGGACTTCGGGCGC 110
 CC |
 CC QY 61 GAGTATGTAAACTCCTGGGCTCTGTGTGTGCTGAGTGCCTGCTACT 111
 CC |
 CC DB 111 GAGTATGTAAACTCCTGGGCTCTGTGTGTGCTGAGTGCCTGCTACT 161
 CC |
 CC
 CC RESULT 2
 CC AA42680
 CC ID AA42680 standard; cDNA; 447 BP.
 CC XX
 CC AC AA42680;
 CC XX
 CC DT 01-FEB-2000 (first entry)
 CC XX
 CC DE Human 5' EST isolated from a cDNA library SEQ ID NO:439.
 CC XX
 CC KM Human: 5' EST; expressed sequence tag; secreted protein; diagnosis;
 CC KM gene therapy; chromosome mapping; upstream regulatory sequence;
 CC KM forensic; location; development; protein synthesis; stability;
 CC KM regulation; identification; ss.
 CC
 CC Homo sapiens.
 CC XX
 CC PN MO9953051-A2.
 CC XX
 CC PD 21-OCT-1999.
 CC XX
 CC PF 09-APR-1999; 99WO-IB00712.
 CC XX
 CC PR 09-APR-1998; 98US-0057719.
 CC PR 28-APR-1998; 98US-0069047.
 CC XX
 CC PA (GEST) GENSET.
 CC XX
 CC PI Dunas Milne Edwards J, Duclert A, Giordano J;
 CC DR WPI: 2000-038446/03.
 CC DR P-PSDB: AAY65066.
 CC XX
 CC Novel secreted protein 5' expressed sequence tag sequences used in
 CC PT diagnostic, forensic, gene therapy, and chromosome mapping procedures
 CC PS Claim 1; Page 402; 837pp; English.
 CC XX
 CC AA42265 to AA43075 represent novel 5' expressed sequence tag (EST)

CC sequences, corresponding to human secreted proteins. AAY64651 to
 CC AAY65438 represent the EST-related proteins corresponding to AA42265 to
 CC AA43052. The 5' ESTs can be used for producing secreted human gene
 CC products. They can be used to identify and isolate 5' untranslated
 CC regions (UTRs) and upstream regulatory regions which control the
 CC location, development stage, rate, and quantity of protein synthesis, as
 CC well as stability of mRNA. The ESTs are also useful as probes for
 CC chromosome mapping, and to obtain full length cDNA clones. The ESTs can
 CC also be used in forensic procedures to identify individuals, or in
 CC diagnostic procedures to identify individuals having genetic diseases
 CC resulting from abnormal gene expression. The products may also be used in
 CC gene therapy protocols. The nucleic acids encoding signal peptides can be
 CC used for directing extracellular secretion of a polypeptide or the
 CC insertion of a polypeptide into a membrane, or importing a polypeptide
 CC into a cell. The proteins encoded by the EST sequences may be useful in
 CC treating a variety of human conditions. Secreted proteins have
 CC therapeutic value, and the identification of new secreted proteins is
 CC valuable. AA42249 to AA42264 and AAY64644 to AAY64650 represent
 CC sequences used in the exemplification of the present invention.
 CC
 CC SQ Sequence 447 BP; 74 A; 105 C; 125 G; 138 T; 5 other;
 CC
 CC Query Match 100.0%; Score 111; DB 21; Length 447;
 CC Best Local Similarity 100.0%; Pred. No. 1.4e-27;
 CC Matches 111; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 CC
 CC QY 1 ATGGGTGATCTTTGCGTGCAGATTCCTTTTCATCTTTGACAGGACTTCGGGCGC 60
 CC |
 CC DB 51 ATGGGTGATCTTTGCGTGCAGATTCCTTTTCATCTTTGACAGGACTTCGGGCGC 110
 CC |
 CC QY 61 GAGTATGTAAACTCCTGGGCTCTGTGTGTGCTGAGTGCCTGCTACT 111
 CC |
 CC DB 111 GAGTATGTAAACTCCTGGGCTCTGTGTGTGCTGAGTGCCTGCTACT 161
 CC |
 CC
 CC RESULT 3
 CC AAF65888/c
 CC ID AAF65888 standard; cDNA; 349 BP.
 CC XX
 CC AC AAF65888;
 CC XX
 CC DT 09-APR-2001 (first entry)
 CC XX
 CC DE Novel human polynucleotide, SEQ ID NO: 1644.
 CC XX
 CC KM Human: cytostatic; gene therapy; colon cancer; prostate cancer;
 CC KM breast cancer; lung cancer; cancer detection; ss.
 CC KM
 CC OS Homo sapiens.
 CC OS
 CC PN WO200102568-A2.
 CC XX
 CC PD 11-JAN-2001.
 CC XX
 CC PF 30-JUN-2000; 2000WO-US18374.
 CC XX
 CC PR 02-JUL-1999; 99US-0142310.
 CC PR 02-JUL-1999; 99US-0142311.
 CC XX
 CC PA (CHIR) CHIRON CORP.
 CC PA (HYSE-) HYSEQ INC.
 CC XX
 CC PI Williams LR, Escobedo J, Innis MA, Garcia PD, Klinger J, Kassam A;
 CC PI Reinhard C, Randazzo F, Kennedy GC, Pot D, Lamson G, Drmanac R;
 CC PI Chtenjakov R, Drmanac S, Dickson M, Iabat I, Leshkowitz D;
 CC PI Kita D, Garcia V, Jones LM, Strache-Crain B;
 CC XX
 CC DR WPI: 2001-091805/10.
 CC XX
 CC Library of polynucleotides for diagnosing a cancerous state of a
 CC PT mammalian cell and detecting cancer, particularly of the colon or
 CC PT prostate, comprises 3351 human polynucleotide sequences -
 CC XX

CC The invention relates to a single exon nucleic acid probe for

CC monitoring and prognosing diseases of the human heart and vascular system

CC congenital heart disease.
CC Note: The sequence data for this patent did not form part of the printed
CC specification, but was obtained in electronic format directly from WIPO
CC at ftp.wipo.int/pub/publsequences.
XX

Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 52.4%; Score 58.2; DB 22; Length 570;
Best Local Similarity 79.3%; Pred. No. 8.2e-10;
Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

24 GGATTCCTTTTCATCTTTCAGGAGCTTCGGGCGGAGATGTAAGTCTGGGCT 83
DB 566 GAATGGATCTCCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 507

84 CTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 110
DB 506 TTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 480

LT 6
ID 11985/c
AAK11985 standard; DNA: 570 BP.

AAK11985;

05-NOV-2001 (first entry)

Human brain expressed single exon probe SEQ ID NO: 11976.

Human; brain expressed exon; gene expression analysis; probe;
microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;
epilepsy; cancer; ss.

Homo sapiens.

MO200157275-A2.

09-AUG-2001.

30-JAN-2001; 2001WO-US00667.

04-FEB-2000; 2000US-0180312.
26-MAY-2000; 2000US-0207456.
30-JUN-2000; 2000US-0608408.
03-AUG-2000; 2000US-0632366.
21-SEP-2000; 2000US-0234687.
27-SEP-2000; 2000US-0236359.
04-OCT-2000; 2000GB-0024263.

(MOLE-) MOLECULAR DYNAMICS INC.

Penn SG, Hanzel DK, Chen W, Rank DR;

WPI; 2001-483446/52.

Single exon nucleic acid probes for analyzing gene expression in human
brains -

Example 4; SEQ ID NO: 11976; 650bp + Sequence Listing; English.

The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC brain. They can be used to measure gene expression in brain cell samples,
CC which may enable the diagnosis and improved treatment of nervous system
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,
CC epilepsy and cancers. The present sequence is one of the probes of the
CC invention.

Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 52.4%; Score 58.2; DB 22; Length 570;
Best Local Similarity 79.3%; Pred. No. 8.2e-10;

Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

24 GGATTCCTTTTCATCTTTCAGGAGCTTCGGGCGGAGATGTAAGTCTGGGCT 83
DB 566 GAATGGATCTCCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 507

84 CTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 110
DB 506 TTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 480

RESULT 7
AAK37688/c
ID AAK37688 standard; DNA: 570 BP.

AAK37688;

06-NOV-2001 (first entry)

Human bone marrow expressed single exon probe SEQ ID NO: 12245.

Human; bone marrow expressed exon; gene expression analysis; probe;
microarray; cancer; leukaemia; lymphoma; myeloma; ss.

Homo sapiens.

MO200157276-A2.

09-AUG-2001.

30-JAN-2001; 2001WO-US00668.

04-FEB-2000; 2000US-0180312.
26-MAY-2000; 2000US-0207456.
30-JUN-2000; 2000US-0608408.
03-AUG-2000; 2000US-0632366.
21-SEP-2000; 2000US-0234687.
27-SEP-2000; 2000US-0236359.
04-OCT-2000; 2000GB-0024263.

(MOLE-) MOLECULAR DYNAMICS INC.

Penn SG, Hanzel DK, Chen W, Rank DR;

WPI; 2001-488900/53.

Human genome-derived single exon nucleic acid probes useful for
analyzing gene expression in human bone marrow -

Example 4; SEQ ID NO: 12245; 658bp + Sequence Listing; English.

The present invention provides a number of single exon nucleic acid
CC probes which are derived from genomic sequences expressed in the human
CC bone marrow. They can be used to measure gene expression in bone marrow
CC samples, which may enable the improved diagnosis and treatment of cancers
CC such as lymphoma, leukaemia and myeloma. The present sequence is one of
CC the probes of the invention.

Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;

Query Match 52.4%; Score 58.2; DB 22; Length 570;
Best Local Similarity 79.3%; Pred. No. 8.2e-10;
Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

24 GGATTCCTTTTCATCTTTCAGGAGCTTCGGGCGGAGATGTAAGTCTGGGCT 83
DB 566 GAATGGATCTCCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 507

84 CTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 110
DB 506 TTGTGTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCTGCT 480

	RESULT 8
A011847/c	ID A0118447 standard; DNA: 570 BP.
XX AC	AA118447;
XX DT	12-OCT-2001 (first entry)
XX XX	Probe #8360 for gene expression analysis in human cervical cell sample.
DE KW	Probe; human; microarray; gene expression; cervical epithelial cell;
XW KW	cervical cancer; ss.
XX OS	Homo sapiens.
XX PN	WO200157278-A2.
XX PD	09-AUG-2001.
XX XX	30-JAN-2001; 2001WO-US00670.
PR PA	04-FEB-2000; 2000US-0180312.
PR PR	26-MAY-2000; 2000US-0207456.
PR PR	30-JUN-2000; 2000US-0608408.
PR PR	03-AUG-2000; 2000US-0632366.
PR PR	21-SEP-2000; 2000US-0234687.
PR PR	27-SEP-2000; 2000US-0236359.
PR PR	04-OCT-2000; 2000GB-0024263.
PA MOLE-	(MOLE-) MOLECULAR DYNAMICS INC.
Penn SG,	Hanzel DK, Chen W, Rank DR; WPI: 2001-488901/53. Human genome-derived single exon nucleic acid probes useful for analyzing gene expression in human cervical epithelial cells - Claim 25; SEQ ID No 8380; 487bp; English.
The present invention relates to human single exon nucleic acid probes (SENP). The present sequence is one such probe. The SENPs are derived from human HeLa cells. The SENPs can be used to produce a single exon microarray, which can be used for measuring human gene expression in a disease derived from human cervical epithelial cells. By measuring gene expression, the probes are therefore useful in grading and/or staging of diseases of the cervix, notably cervical cancer. Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO at ftp.wipo.int/pub/published_pcl_sequences.	
Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;	
Query Match 52.4%; Score 58.2; DB 22; Length 570; Best Local Similarity 79.3%; Pred. No. 8.2e+10; Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;	
OY 24 GGATTCCTTTTATTGAGCAGACCTTGCGCCGGAGTAGTAATAAACCCTGGGTCT 83 DB 566 GAATGCATCTCTGCTCTGCTCTGGCATCTCTTGCGCTGGAATATGTAATAATCTGGGTCT 507 OY 84 CTGTGTGTCCTGAGTGGCTGCTTAC 110 DB 506 TTGTGTGTGCCCTGAGTGGCGCCTCTGC 480 RESULTS 9 AAI43563/c ID AAI43563 standard; DNA: 570 BP. AC AA143563; DT 17-OCT-2001 (first entry) XX	

XX	Probe #12249 used to measure gene expression in human placenta sample.
XX	
KW	Probe; microarray; human; placenta; antenatal diagnosis;
KW	genetic disorder; ss.
XX	
OS	Homo sapiens.
XX	
FN	WO200157272-A2.
XX	
PD	09-AUG-2001.
XX	
PF	30-JAN-2001; 2001WO-US00663.
XX	
PR	04-FEB-2000; 2000US-0180312.
PR	26-MAY-2000; 2000US-0207456.
PR	30-JUN-2000; 2000US-0608408.
PR	03-AUG-2000; 2000US-0633366.
PR	21-SEP-2000; 2000US-0234687.
PR	27-SEP-2000; 2000US-0236359.
PR	04-OCT-2000; 2000GB-0024263.
PA	(MOLE-) MOLECULAR DYNAMICS INC.
XX	
PI	Penn SG, Hanzel DK, Chen W, Rank DR;
XX	
DR	WPI; 2001-488897/53.
XX	
PT	Human genome-derived single exon nucleic acid probes useful for analyzing gene expression in human placenta -
XX	
PS	Claim 25; SEQ ID No 12249; 654pp; English.
XX	
CC	The present invention relates to single exon nucleic acid probes (SENPs).
CC	The present sequence is one such probe. The probes are useful for
CC	producing a microarray for predicting, measuring and displaying gene
CC	expression in samples derived from human placenta. The probes are useful
CC	for antenatal diagnosis of human genetic disorders.
XX	
SQ	Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
	Query Match 52.4%; Score 58.2; DB 22; Length 570;
	Best Local Similarity 79.3%; Pred. No. 8.2e-10;
	Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0.
OY	24 GCATCTTTTTCATCTTCACAGGACTTCGGCGCGGAGTAATGAACCTCCTGCCTCT 83 Db 566 GAATGATCTCTCGCTTGCTGCTGCGATTCCTTGCGCTGGAGTATGTAAATTCTCGGCTCT 507
OY	84 CTGTGTGCTCCTGAGTGGCTGCTTAC 110 Db 506 TTGTGTGCTCGTAGTGCGCGCTCTGC 480
RESULT 10	
ABS11680/C	
ID	ABS11680 standard; DNA; 570 BP.
XX	
AC	ABS11680;
XX	
DT	19-AUG-2002 (first entry)
XX	
DE	Human genome-derived single exon probe from lung SEQ ID No 11671.
XX	
KW	Human; ds; single exon probe; asthma; lung cancer; COPD; ILD;
KW	chronic obstructive pulmonary disease; interstitial lung disease;
KW	familial idiopathic pulmonary fibrosis; neurofibromatosis;
KW	tuberosclerosis; Gaucher's disease; Niemann-Pick disease;
KW	Hernansky-Rudrak syndrome; sarcoidosis; pulmonary haemosiderosis;
KW	pulmonary histiocytosis; lymphangioleiomyomatosis; Karagener syndrome;
KW	pulmonary alveolar proteinosis; fibrocystic pulmonary dysplasia;
KW	primary ciliary dyskinesia; pulmonary hypertension;
KW	hyaline membrane disease.

XX Homo sapiens.
 OS WO200186003-A2.
 PN 15-NOV-2001.
 PD 30-JAN-2001; 2001WO-US00665.
 PF 04-FEB-2000; 2000US-180312P.
 PR 26-MAY-2000; 2000US-207456P.
 PR 30-JUN-2000; 2000US-0608408.
 PR 03-AUG-2000; 2000US-0632366.
 PR 21-SEP-2000; 2000US-234687P.
 PR 27-SEP-2000; 2000US-236359P.
 PR 04-OCT-2000; 2000GB-0024263.
 XX (MOLE-) MOLECULAR DYNAMICS INC.
 PA Penn SG, Hanzel DK, Chen W, Rank DR;
 XX
 WPI: 2002-114183/15.
 PT Spatially-addressable set of single exon nucleic acid probes, used to
 PT measure gene expression in human lung samples -
 PS
 PS Claim 1: SEQ ID No 11671; 634pp; English.
 XX
 XX The invention relates to a spatially-addressable set of single exon
 CC nucleic acid probes for measuring gene expression in a sample derived
 CC from human lung comprising single exon nucleic acid probes having one of
 CC 12614 nucleic acid sequences mentioned in the specification, or their
 CC complements or the 12387 open reading frames derived from the 12614
 CC probes. Also included are a microarray comprising the novel set of
 CC probes; the novel set of probes which hybridize at high stringency to a
 CC nucleic acid expressed in the human lung; measuring gene expression in a
 CC sample derived from human lung, comprising (a) contacting the array with
 CC a collection of detectably labeled nucleic acids derived from human lung
 CC mRNA, and (b) measuring the label detectably bound to each probe of
 CC the array; identifying exons in a eukaryotic genome, comprising
 CC (a) algorithmically predicting at least one exon from genomic sequences
 CC of the eukaryote; and (b) detecting specific hybridisation of detectably
 CC labeled nucleic acids from eukaryote lung mRNA, to a single exon probe,
 CC having a fragment identical to the predicted exon, the probe is included
 CC in the above mentioned microarray; assigning exons to a single gene,
 CC comprising (a) identifying exons from genomic sequence by the method
 CC above and (b) measuring the expression of each of the exons in several
 CC tissues and/or cell types using hybridisation to a single exon
 CC microarrays having a probe with the exon, where a common pattern of
 CC expression of the exons in the tissues and/or cell types indicates that
 CC the exons should be assigned to a single gene; a peptide comprising one
 CC of 12011 sequences, mentioned in the specification, or encoded by the
 CC probes/open reading frames (ORF). The probes are used for gene
 CC expression analysis, and for identifying exons in a gene, particularly
 CC using human lung derived mRNA and for the study of lung diseases
 CC such as asthma, lung cancer, chronic obstructive pulmonary disease
 CC (COPD), interstitial lung disease (ILD), familial idiopathic pulmonary
 CC fibrosis, neurofibromatosis, tuberous sclerosis, Gaucher's disease,
 CC Niemann-Pick disease, Hermansky-Pudlak syndrome, sarcoidosis, pulmonary
 CC haemosiderosis, pulmonary histiocytosis, lymphangioleiomyomatosis,
 CC pulmonary alveolar proteinosis, Karagenen syndrome, fibrocystic
 CC pulmonary dysplasia, primary ciliary dyskinesia, pulmonary hypertension
 CC and hyaline membrane disease. The present sequence is a single exon
 CC probe of the invention.
 CC Note: The sequence data for this patent did not form part
 CC of the printed specification, but was obtained in electronic
 CC format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences.
 XX
 XX Sequence 570 BP; 169 A; 154 C; 156 G; 91 T; 0 other;
 Query Match 52.4%; Score 58.2; DB 24; Length 570;
 Best Local Similarity 79.3%; Pred. No. 8.2e-10;

Matches 69; Conservative 0; Mismatches 18; Indels 0; Gaps 0;
 QY 24 GGATTCCTTTTTCATCTTTGACGAGACTTCTGGGCGCGAGTATGTAACCTCTGGGCT 83
 Db 566 GAATGATCTCCCTGCTGCTGGATCTCTGGCTGAGATGTAATCTCTGGGCT 507
 QY 84 CTGTGTGTGCTGAGTGGCTGCTTAC 110
 Db 506 TTGTGTGTGCTGAGTGGCTGCTTAC 480
 RESULT 11
 ABR83461/C
 ID ABR83461 standard; cDNA: 128600 BP.
 AC ABR83461;
 XX
 XX 14-AUG-2002 (first entry)
 DT
 DT Human cDNA differentially expressed in granulocytic cells #32.
 DE
 XX Human; ss; granulocytic cell; DNA chip; bacterial infection;
 KW viral infection; parasitic infection; protozoal infection;
 KW fungal infection; sterile inflammatory disease; psoriasis;
 KW rheumatoid arthritis; glomerulonephritis; asthma; thrombosis;
 KW cardiac reperfusion injury; renal reperfusion injury; ARDS;
 KW adult respiratory distress syndrome; inflammatory bowel disease;
 KW Crohn's disease; ulcerative colitis; periodontal disease;
 KW granulocyte activation; chronic inflammation; allergy.
 KW
 XX Homo sapiens.
 OS
 OS WO200228999-A2.
 PN 11-APR-2002.
 PD
 PD 03-OCT-2001; 2001WO-US30821.
 PF
 PF 03-OCT-2000; 2000US-237189P.
 PR
 PR (GENE-) GENE LOGIC INC.
 PA Beazer-Barclay Y, Weissman SM, Yamaga S, Vockley J;
 PI
 PI WPI: 2002-435328/46.
 DR
 DR
 PT Detecting granulocyte activation by detecting differential expression
 PT of genes associated with granulocyte activation, which serves as
 PT diagnostic markers that is useful for monitoring disease states and
 PT drug toxicity -
 PS
 PS Claim 1: SEQ ID No 32; 114pp; English.
 XX
 XX The invention relates to detecting (M1) granulocyte (GC) activation
 CC (GCA), by detecting the level of expression of gene(s) (Gs) identified by
 CC DNA chip analysis as given in the specification, and comparing
 CC the expression level to an expression level in an unactivated
 CC GC, where differential expression of Gs is indicative of GCA.
 CC Also included are modulating (M2) Gs by contacting GC with an agent
 CC that alters the expression of at least one gene in Gs; (2) screening (M3)
 CC for an agent capable of modulating GCA or an inflammation (especially
 CC chronic) in a tissue, an allergic response in a subject, exposure of a
 CC subject to a pathogen or sterile inflammatory disease using the
 CC gene expression profile; (3) detecting (M4) an inflammation (especially
 CC chronic) in a tissue, an allergic response in a subject, exposure of a
 CC subject to a pathogen or sterile inflammatory disease, by detecting the
 CC level of expression in a sample of the tissue of gene(s) from Gs, where
 CC the level of expression of the gene is indicative of inflammation;
 CC (4) treating (M5) an inflammation (especially chronic) or in a tissue,
 CC an allergic response in a subject, exposure of a subject to a pathogen
 CC or sterile inflammatory disease, by contacting a tissue having
 CC inflammation with an agent that modulates the expression of gene(s)
 CC from Gs in the tissue. M1 is useful for detecting GCA; M2 is useful for

modulating GA: M3 is useful for screening an agent capable of modulating
CCA preferably in an inflammation in a tissue; M4 is useful for
detecting an inflammation (especially chronic) in a tissue, an allergic
response in a subject, exposure of a subject to a pathogen or sterile
inflammatory disease (e.g. psoriasis, rheumatoid arthritis, renal
glomerulonephritis, asthma, thrombosis, cardiac reperfusion injury, renal
reflexion injury, ARDS, adult respiratory distress syndrome,
inflammatory bowel disease, Crohn's disease, ulcerative colitis,
periodontal disease; also bacterial infection, viral infection,
parasitic infection, protozoal infection, fungal infection and M5 is
useful for treating one of the above conditions. The present
sequence represents a gene differentially expressed in granulocytes.
Note: The sequence data for this patent did not form part
of the printed specification, but was obtained in electronic
format directly from WIPO at
ftp://wipo.int/pub/published_pct_sequences.

Sequence 128600 BP; 36139 A; 24970 C; 26316 G; 41175 T; 0 other;

Query Match 50.8%; Score 56.4; DB 24; Length 128600;

Best Local Similarity 70.8%; Pred. No. 1.6e-08; Mismatches 31; Indels 0; Gaps 0;

Matches 75; Conservative 0; Mismatches 31; Indels 0; Gaps 0;

DB 126322 ATGTGTGACCTCCACCTTCACTTCAATTCAGACGACATTTGTGGGGCTCTGGGGCTG 126263

QY 61 GAGTATGTAACACTCTGGGCTCTGTGTGCTGAGTGGCTGCT 106

DB 126262 GAGTATGTAACACTCTGGGCTCTGTGTGCTGAGTGGCTGCT 126217

RESULT 12

AAC68089/c AAC68089 standard; cDNA: 1982 BP.

AC AAC68089;

DT 20-FEB-2001 (first entry)

XX Human secreted protein cDNA sequence #9.

XX Cytostatic; immunosuppressive; nootropic; neuroprotective; antiviral;
XX allergic; hepatotropic; antidiabetic; antiinflammatory; antitumor;
XX antineoplastic; anticonvulsant; antibacterial; antifungal; antiparasitic;
XX cardiant; gene therapy; cancer; immune disorder; cardiovascular disorder;
XX neurological disease; infection; human; secreted protein; ss.

XX Homo sapiens.

PN WO200058335-A1.

XX 05-OCT-2000.

PF 22-MAR-2000; 2000WO-US07534.

PR 26-MAR-1999; 99US-0126598.

PR 22-DEC-1999; 99US-0171504.

XX (HUMA-) HUMAN GENOME SCI INC.

XX (ROSE/) ROSEN C A.

XX Rosen CA, Ruben SM, Komatsoulis G;

XX WPI: 2000-611702/58.

XX P-PSDB: AAB37336.

XX Nucleic acids encoding human secreted proteins, used to treat, prevent,
XX ameliorate or diagnose conditions such as cancer, and autoimmune
XX diseases e.g. arthritis -
XX Claim 1; Pages 321-322; 387pp; English.

The invention relates to the isolation of genes AAC68081-C68127 encoding
CC 47 human secreted proteins AAB37348-B37394. The genes can be used to
CC generate fusion proteins by linking to the gene for the human
CC immunoglobulin G Fc portion (AAC68072) for increasing the stability of
CC the fusion protein as compared to the human protein only. The genes and
CC proteins are useful for preventing, ameliorating or treating medical
CC conditions, e.g. by protein or gene therapy. The genes are isolated
CC from a range of human tissues disclosed in the specification. The
CC nucleic acids, proteins, antibodies and (ant)agonists are useful in
CC the diagnosis, treatment and prevention of: (a) cancer, e.g. breast
CC and ovarian cancer, and other cancers of the adrenal gland, bone, bone
CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;
CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune
CC hemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative
CC colitis; (c) cardiovascular disorders such as myocardial ischemias; (d)
CC wound healing; (e) neurological diseases e.g. cerebral anoxia and
CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal
CC and parasitic infections.

Sequence 1982 BP; 549 A; 513 C; 471 G; 448 T; 1 other;

Query Match 49.5%; Score 55; DB 21; Length 1982;

Best Local Similarity 70.9%; Pred. No. 1.4e-08; Mismatches 30; Indels 0; Gaps 0;

Matches 73; Conservative 0; Mismatches 30; Indels 0; Gaps 0;

QY 9 ATCTTTGACCTTGACGATTTCTTTTCATCTTTCAGACGACTCTGGGCGGAGATATGT 68

DB 1163 ATCTCCCGCTTTGACGAGTGTGCACTTCTTGTGGGAAACCAAGCGCGGAGATATGT 1104

QY 69 AAAATCTGCGGCTCTGTGTGCTGAGTGGCTGCTCTACT 111

DB 1103 AAAATCTGCGGCTCTGTGTGCTGAGTGGCTGCTCTACT 1061

RESULT 13

AAH98980 ID AAH98980 standard; cDNA: 609 BP.

AC AAH98980;

DT 12-OCT-2001 (first entry)

XX Human EST-derived coding sequence SEQ ID NO: 837.

XX Human; sheep; pig; cow; fruit fly; yeast; hamster; macaque; horse;
XX tomato; monkey; dog; sea urchin; expressed sequence tag; EST;
XX diagnostics; forensic test; gene mapping; genetic disorder;
XX biodiversity; gene therapy; nutrition; ss.

XX Homo sapiens.

PN WO200154477-A2.

XX 02-AUG-2001.

PF 25-JAN-2001; 2001WO-US02687.

PR 25-JAN-2000; 2000US-0491404.

PR 17-JUL-2000; 2000US-0617746.

PR 03-AUG-2000; 2000US-0631451.

PR 15-SEP-2000; 2000US-0653870.

XX (HYSE-) HYSEQ INC.

XX Tang YT, Liu C, Zhou P, Qian XB, Wang Z, Chen R, Asundi V;

XX Cao Y, Drmanac RA, Zhang J, Weinman T;

XX WPI: 2001-476164/51.

XX P-PSDB: AAM24321.

XX Isolated polypeptide for treatment of diseases, diagnostics, raising
XX antibodies and research use -

GenCore version 5.1.4-p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 01:47:50 ; Search time 1042 Seconds

(without alignments)
1725.239 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggtgatcttgcctt.....gcttgagtgctgctact 111

Scoring table: IDENTIFY_NUC
Gapop 10.0 ; Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

1 number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%

Listing first 45 summaries

Database :

BST:*

- 1: em_estba:*
- 2: em_esthum:*
- 3: em_estlin:*
- 4: em_estlmu:*
- 5: em_estlov:*
- 6: em_estlpl:*
- 7: em_estro:*
- 8: em_hlc:*
- 9: gb_estl:*
- 10: gb_estl2:*
- 11: gb_hlc:*
- 12: gb_estl3:*
- 13: gb_estl4:*
- 14: gb_estl5:*
- 15: em_estlun:*
- 16: em_estlom:*
- 17: gb_gss:*
- 18: em_gss_hum:*
- 19: em_gss_inv:*
- 20: em_gss_pln:*
- 21: em_gss_vrt:*
- 22: em_gss_fun:*
- 23: em_gss_mam:*
- 24: em_gss_mus:*
- 25: em_gss_other:*
- 26: em_gss_pro:*
- 27: em_gss_rod:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	111	100.0	470 17	A0770688	A0770688 HS_5368_B
2	99.4	89.5	525 17	A0165256	A0165256 HS_3025_B
3	63.4	57.1	519 17	A0186743	A0186743 HS_3109_B
4	63	56.8	416 17	A0182486	A0182486 HS_3077_A
5	63	56.8	867 12	BG675164	BG675164 602621440
6	61.4	55.3	516 17	A0457001	A0457001 HS_5151_A

C 7	58.8	53.0	529 17	A0881246	A0881246 HS_5137_B
C 8	58.4	52.6	597 17	A0506984	A0506984 RPT-11-3
9	58.2	52.4	416 17	B75884	B75884 RPT-11-12L1
C 10	58.2	52.4	530 17	A0193128	A0193128 HS_3060_B
11	57.8	52.1	448 17	A0702903	A0702903 HS_5452_B
12	57.8	52.1	769 17	A0899390	A0899390 HS_5234_A
13	57.6	51.9	413 17	A0266658	A0266658 RPT-11-73
14	57.2	51.5	453 17	A0442274	A0442274 HS_5137_A
15	57.2	51.5	635 17	A0390599	A0390599 HS_5137_A
C 16	56.8	51.2	519 17	A0399984	A0399984 HS_3106_A
17	56.6	51.0	491 17	A0186162	A0186162 HS_3077_A
18	56.6	51.0	628 17	A0237815	A0237815 RPT-11-70
C 19	56.6	51.0	864 17	A0739814	A0739814 HS_5505_A
20	56.6	50.5	546 17	A0637256	A0637256 RPT-11-4
21	56.6	50.1	513 17	A0207172	A0207172 HS_3239_B
22	55.6	50.1	513 17	A0455447	A0455447 HS_5153_A
23	55.6	50.1	513 17	A0765592	A0765592 HS_5348_B
24	55.6	50.1	541 17	A0451696	A0451696 RPT-11-3
25	55.6	50.1	546 17	A0427698	A0427698 CITBI-E1-
26	55.6	50.1	732 17	A0306035	A0306035 RPT-11-3
C 27	55.4	49.9	580 17	A0532835	A0532835 RPT-11-3
28	55.4	49.7	695 17	AG179297	AG179297 Pan trogl
C 29	55.2	49.5	423 17	A0564722	A0564722 HS_5361_A
30	55	49.5	482 17	A0320567	A0320567 RPT-11-99
C 31	55	49.5	549 17	A0540868	A0540868 RPT-11-3
32	55	49.5	551 17	A0569689	A0569689 HS_5333_B
33	55	49.5	563 17	A0420187	A0420187 RPT-11-1
C 34	55	49.5	580 17	A0316169	A0316169 RPT-11-10
35	55	49.5	634 17	AG160901	AG160901 Pan trogl
36	55	49.5	659 17	AG151043	AG151043 Pan trogl
37	55	49.5	723 17	A0386439	A0386439 RPT-11-15
C 38	54.8	49.4	483 17	A0508480	A0508480 RPT-11-2
39	54.6	49.2	483 17	A0668395	A0668395 HS_5414_A
40	54.6	49.2	653 17	AG143347	AG143347 Pan trogl
C 41	54.6	49.2	683 17	AG091225	AG091225 Pan trogl
42	54.6	49.2	503 17	AQ155611	AQ155611 HS_3124_A
43	54.4	49.0	708 17	AG092466	AG092466 Pan trogl
C 44	54.4	49.0	672 17	AG051939	AG051939 Pan trogl
45	54.2	48.8			

ALIGNMENTS

RESULT 1
LOCUS A0770688
DEFINITION HS_5368_B2.C08.SP6E RPT-11 Human Male BAC library Homo sapiens
ACCESSION A0770688
VERSION A0770688.1 GI:5648804
KEYWORDS GSS.
SOURCE human.
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homiidae; Homo.
AUTHORS Mahairas,G.G., Wallace,J.C., Smith,K., Swartzell,S., Holzman,T., Keller,A., Shaker,R., Furlong,J., Young,J., Zhao,S., Adams,M.D. and Hood,L.
Sequence-tagged connectors: A sequence approach to mapping and scanning the human genome
Proc. Natl. Acad. Sci. U.S.A. 96 (17), 9739-9744 (1999)
CONTACT: Mahairas GG, Wallace JC, Hood L
High Throughput Sequencing Center
University of Washington
401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618
Fax: (206) 616-3887
Email: jwallace@u.washington.edu
Clones are derived from the human BAC library RPT-11. For BAC library availability, please contact Pieter de Jong

[illegible]

401 Queen Anne Avenue North, Seattle, WA 98109, USA
Tel: (206) 616-3618

Department of Eukaryotic Genomics
The Institute for Genomic Research
9712 Medical Center Dr., Rockville, MD 20850, USA
Tel: 301 838 0200
Fax: 301 838 0208
Email: mdadam@tigr.org
Clones are derived from the human BAC library RPc1-11. For BAC library availability, please contact Pieter de Jong (pieter@dejong.med.buffalo.edu). Clones may be purchased from BACPAC Resources (<http://bacpac.med.buffalo.edu/ordering>) or from Research Genetics (info@resgen.com). BAC end search page: http://www.tigr.org/tldb/hungen/bac_end_search/bac_end_search.html
Seq primer: SP6
Class: BAC ends.

	Matches	68;	Conservative	0;	Mismatches	18;	Indels	0;	Gaps	0;
QY	25	GATTCTTTTTCATCTTTGCGAGGACTTCTGCGCGCGAGATATGTAATAACTCCTGGGCTC	84							
Db	114	GATGGTTCTTCGCTGCTGCTGATCCAGGCGCAGAGTATGCAAAATTCCTGGGCTC	173							
QY	85	TGTGTGTGCTGAGTGGCTGCTCTAC	110							
Db	174	TGTGTGTGCTGAGTGGCTGCTCTGC	199							

Search completed: April 25, 2003, 02:28:38
 Job time : 1045 secs

GenCore version 5.1.4-p5_4578
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 01:55:25 ; Search time 39 Seconds
(without alignments)
872.849 Million cell updates/sec

Title: US-09-513-999C-3792_COPY_51_161

Perfect score: 111

Sequence: 1 atgggtgagctcttgcctt.....gcctgagtgctgcttact 111

Scoring table: IDENTITY NUC

Gapop 10.0, Gapept 1.0

Searched: 441362 seqs, 15338381 residues

1 number of hits satisfying chosen parameters: 882724

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents.NA.*

1: /cgn2_6/ptodata/1/ina/5A.COMB.seq.*
2: /cgn2_6/ptodata/1/ina/5B.COMB.seq.*
3: /cgn2_6/ptodata/1/ina/6A.COMB.seq.*
4: /cgn2_6/ptodata/1/ina/6B.COMB.seq.*
5: /cgn2_6/ptodata/1/ina/PCUTS.COMB.seq.*
6: /cgn2_6/ptodata/1/ina/backfile1.seq.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	27.6	24.9	80246	4	US-09-078-294-4
2	27.6	24.9	80595	4	US-09-078-294-3
3	26.2	23.6	175	1	US-08-222-177A-4
4	26.2	23.6	1146	2	US-08-666-367B-4
5	26.2	23.6	1146	4	US-09-143-438-4
6	26.2	23.6	13104	4	US-08-961-527-34
7	26.2	23.4	430	4	US-09-397-787-254
8	25.4	22.9	11492	4	US-08-991-840A-1
9	24.8	22.3	1371	2	US-08-428-713-1
10	24.8	22.3	1371	3	US-08-904-179-1
11	24.8	22.3	1374	2	US-08-428-713-9
12	24.8	22.3	1374	3	US-08-904-179-9
13	24.6	22.2	882	4	US-09-556-877-136
14	24.6	22.2	882	4	US-09-620-412C-136
15	24.6	22.2	2407	4	US-09-370-807-7
16	24.6	22.2	2407	4	US-09-921-259-7
17	24.6	22.2	2511	4	US-09-422-869-19
18	24.4	22.0	361	4	US-09-385-982-26
19	24.4	22.0	1506	4	US-08-206-790A-22
20	24.4	22.0	1506	5	PCT-US95-02843-22
21	24.2	21.8	2347	1	US-08-145-681-3
22	24.2	21.8	2347	1	US-08-453-703-3
23	24.2	21.8	2347	2	US-08-456-106-3
24	24.2	21.8	2347	3	US-08-456-108-3
25	24.2	21.8	2347	4	US-09-265-577-3
26	24	21.6	528	4	US-09-134-001C-2094
27	24	21.6	4088	2	US-08-317-310A-1

28	24	21.6	4088	5	PCT-US95-13041-1	Sequence 1, Appli
29	24	21.6	19446	4	US-08-961-527-51	Sequence 51, Appl
30	23.8	21.4	3831	4	US-08-961-527-291	Sequence 291, App
31	23.8	21.4	7542	4	US-09-734-030-3	Sequence 3, Appli
32	23.8	21.4	13158	2	US-08-687-080-105	Sequence 105, App
33	23.8	21.4	35100	1	US-08-306-691B-19	Sequence 19, Appl
34	23.8	21.4	35100	3	PCT-US93-06251-19	Sequence 19, Appl
35	23.8	21.4	87350	3	US-08-781-891-79	Sequence 79, Appl
36	23.8	21.4	87543	4	US-09-791-211-3	Sequence 3, Appli
37	23.6	21.3	15567	4	US-09-627-376-3	Sequence 3, Appli
38	23.4	21.1	560	3	US-09-059-369-18	Sequence 18, Appl
39	23.4	21.1	598	4	US-08-998-416-1076	Sequence 1076, Ap
40	23.4	21.1	694	4	US-08-998-416-710	Sequence 710, App
41	23.4	21.1	1477	4	US-09-123-030-9	Sequence 9, Appli
42	23.4	21.1	2090	3	US-09-059-369-1	Sequence 1, Appli
43	23.4	21.1	3093	1	US-08-252-966B-19	Sequence 19, Appli
44	23.4	21.1	5253	2	US-08-290-731C-3	Sequence 3, Appli
45	23.2	20.9	401	4	US-09-221-296-40	Sequence 40, Appli

ALIGNMENTS

```
RESULT 1
US-09-078-294-4
; Sequence 4, Application US/09078294
; Patent No. 6265211
; GENERAL INFORMATION:
; APPLICANT: Choo, Kong-Hong Andy
; APPLICANT: Du Sart, Desiree
; APPLICANT: Cancilla, Michael R.
; TITLE OF INVENTION: A NOVEL NUCLEIC ACID MOLECULE
; FILE REFERENCE: Davies Col
; CURRENT APPLICATION NUMBER: US/09/078, 294
; CURRENT FILING DATE: 1998-05-13
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 4
; LENGTH: 80246
; TYPE: DNA
; ORGANISM: Nucleotide sequence of NC-contlig
US-09-078-294-4

Query Match          24.9%; Score 27.6; DB 4; Length 80246;
Best Local Similarity 58.5%; Pred. No. 5.3;
Matches 48; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

QY 23 AGCATTTCTTTTATCTTTCAGGAGCTTGGGCCGAGATGTAACACTCTGGGTC 82
    || |||| |||| |||| |||| |||| |||| |||| |||| |||| ||||
Db 6333 AGTTTCTGTGTCAACTGACGTGGCCATGGATGTCAGATATGTAATTAACAGTATT 6392
QY 83 TCTGTGTGCTGCTGAGTGGCTG 104
    |||| |||| |||| |||| |||| |||| |||| |||| |||| ||||
Db 6393 TCTGGGTGTTTCTGTGAGGGTG 6414

RESULT 2
US-09-078-294-3
; Sequence 3, Application US/09078294
; Patent No. 6265211
; GENERAL INFORMATION:
; APPLICANT: Choo, Kong-Hong Andy
; APPLICANT: Du Sart, Desiree
; APPLICANT: Cancilla, Michael R.
; TITLE OF INVENTION: A NOVEL NUCLEIC ACID MOLECULE
; FILE REFERENCE: Davies Col
; CURRENT APPLICATION NUMBER: US/09/078, 294
; CURRENT FILING DATE: 1998-05-13
; NUMBER OF SEQ ID NOS: 29
; SOFTWARE: Patentln Ver. 2.0
; SEQ ID NO 3
; LENGTH: 80595
; TYPE: DNA
```


TOPOLOGY: Linear
US-08-961-527-34

Query Match 23.6%; Score 26.2; DB 4; Length 13104;
Best Local Similarity 67.3%; Pred. No. 8.6;
Matches 37; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

QY 10 TCTTTGGCTTGCAGGATCTTTTCATCTTTGCAGGACTTGGGCGGAGT 64
DB 9341 TCTATTGCTTGGGGGCTTCTTGGGCACTTTTGTAGGAGATTAAAGCACGCT 9287

RESULT 7

US-09-397-787-254/C
Sequence 254; Application US/09397787
Patent No. 6468758
GENERAL INFORMATION:
APPLICANT: Benson, Darin R.
APPLICANT: Lodes, Michael J.
APPLICANT: Mitcham, Jennifer L.
APPLICANT: King, Gordon E.
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR OVARIAN
TITLE OF INVENTION: CANCER THERAPY AND DIAGNOSIS
FILE REFERENCE: 210121.466C2
CURRENT APPLICATION NUMBER: US/09/397.787
CURRENT FILING DATE: 1999-09-16
NUMBER OF SEQ ID NOS: 334
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 254
LENGTH: 430
TYPE: DNA
ORGANISM: Homo sapien
FEATURE:
NAME/KEY: misc_feature
LOCATION: (1)...(430)
OTHER INFORMATION: n = A,T,C or G
US-09-397-787-254

Query Match 23.4%; Score 26; DB 4; Length 430;
Best Local Similarity 70.0%; Pred. No. 2.9;
Matches 35; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

QY 62 AGATGTAACCTCGGCTCTGTGTGTCCTGAGTGGCTGCTACT 111
DB 402 AGATGTAACCTCGGCTCTGTGTGTCCTGAGTGGCTGCTACT 353

RESULT 8

US-991-840A-1/C
Sequence 1, Application US/08991840A
Patent No. 6261570

GENERAL INFORMATION:
APPLICANT: Michael D. Parker
APPLICANT: Jonathan F. Smith
APPLICANT: Bruce Crise
APPLICANT: Mark Steve Oberste
APPLICANT: Shannon Schmura
TITLE OF INVENTION: Live Attenuated Virus Vaccines for Eastern Equine Encephalitis
NUMBER OF SEQUENCES: 29
CORRESPONDENCE ADDRESS:
ADDRESSEE: Charles H. Harris
STREET: USA MRC - MRC-JA
CITY: FORT DETRICK, FREDERICK
STATE: MARYLAND
COUNTRY: USA
ZIP: 21702-5012

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: Apple Macintosh
OPERATING SYSTEM: Macintosh 7.5
SOFTWARE: Microsoft Word 6.0
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/991.840A

FILING DATE: December 16, 1997
CLASSIFICATION: 424
PRIOR APPLICATION DATA:
APPLICATION NUMBER: Provisional Application 60/047162,
FILING DATE: May 20, 1997
APPLICATION NUMBER: Provisional Application 60/053,652
FILING DATE: July 24, 1997

ATTORNEY/AGENT INFORMATION:
NAME: Charles H. Harris
REGISTRATION NUMBER: 34,616
REFERENCE/DOCKET NUMBER: 003/058/SAP RIID 96-01
TELECOMMUNICATION INFORMATION:
TELEPHONE: (301) 619-2065
TELEFAX: (301) 619-5034
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 11492 base pairs
TYPE: Nucleic acid
STRANDEDNESS: Double

TOPOLOGY: Linear
FEATURE: OTHER INFORMATION: N at all occurrences is = unknown.
FEATURE: OTHER INFORMATION: K at all
FEATURE: occurrences is = G or T
US-08-991-840A-1

Query Match 22.9%; Score 25.4; DB 4; Length 11492;
Best Local Similarity 64.4%; Pred. No. 16;
Matches 38; Conservative 0; Mismatches 21; Indels 0; Gaps 0;

QY 12 TTTTGGCTTGCAGGATCTTTTCATCTTTGCAGGACTTGGGCGGAGTATGTA 70
DB 7692 TTTGCTTGGGAGGACTTCTTCTTCTTCTTGGGCGGAGCTGGCGGAGTATGTA 7634

RESULT 9

US-08-428-713-1/C
Sequence 1, Application US/08428713
Patent No. 5866541

GENERAL INFORMATION:
APPLICANT: Hook, Magnus
APPLICANT: LINDBERG, Kjell Martin
APPLICANT: LINDGREN, Per-Eric
APPLICANT: SIGNAS, Lars Christer
TITLE OF INVENTION: FIBRONECTIN BINDING PROTEIN
NUMBER OF SEQUENCES: 10
CORRESPONDENCE ADDRESS:
ADDRESSEE: Burns, Doane, Swecker & Mathis
STREET: P.O. Box 1404
CITY: Alexandria
STATE: Virginia
COUNTRY: United States
ZIP: 22313-1404

COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/428,713
FILING DATE: 25-APR-1995
CLASSIFICATION: 514

PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/125,222
FILING DATE: 23-SEP-1993
ATTORNEY/AGENT INFORMATION:
NAME: Rea, Teresa Stanek
REGISTRATION NUMBER: 30,427
REFERENCE/DOCKET NUMBER: 012885-074
TELECOMMUNICATION INFORMATION:
TELEPHONE: (703) 836-6620
TELEFAX: (703) 836-2021
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:


```

; APPLICANT: HOOK, Magnus
; APPLICANT: LINDERG, Kjell Martin
; APPLICANT: LINDERG, Per-Eric
; APPLICANT: SIGRAN, Lars Christer
; TITLE OF INVENTION: FIBRONECTIN BINDING PROTEIN
; NUMBER OF SEQUENCES: 10
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Burns, Doane, Swecker & Mathis
; STREET: P.O. Box 1404
; CITY: Alexandria
; STATE: Virginia
; COUNTRY: United States
; ZIP: 22133-1404
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patentln Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/904,179
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/428,713
; FILING DATE: 25-APR-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Rea, Teresa Stanek
; REGISTRATION NUMBER: 30,427
; REFERENCE/DOCKET NUMBER: 012885-074
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 836-6620
; TELEFAX: (703) 836-2021
; INFORMATION FOR SEQ ID NO: 9:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1374 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..1164
; US-08-904-179-9

Query Match          22.3%; Score 24.8; DB 3; Length 1374;
Best Local Similarity 56.0%; Pred. No. 12;
Matches 47; Conservative 0; Mismatches 37; Indels 0; Gaps 0;

13 TTGGCTTGACGAGATCTTTTCATCTTTCGACGAGACTTCGGGCGCGAGTAGTAATAA 72
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 322 TCTGACCCACCAAGATTATTTCTCCCTGTGGACCTTCGTGTGATGATGCTGTGAA 263
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

OY 73 CTCCTGGCTCTCTGTGTGCTG 96
   ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

Db 262 CTGAAGTTCTGACGGATATCAGTG 239

RESULT 13
US-09-556-877-136/c
; Sequence 136, Application US/09556877
; Patent No. 6432816
; GENERAL INFORMATION:
; APPLICANT: Probst, Peter
; APPLICANT: Bhatia, Ajay
; APPLICANT: Skeiky, Yasir
; APPLICANT: Fling, Steve
; APPLICANT: Watsonneuve, Jeff
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATMENT AND
; FILE REFERENCE: 210121.469C5
; TITLE OF INVENTION: DIAGNOSIS OF CHLAMYDIAL INFECTION
; CURRENT APPLICATION NUMBER: US/09/556,877
; CURRENT FILING DATE: 2000-04-19
; NUMBER OF SEQ ID NOS: 305
```

```

; SOFTWARE: FastSeq for Windows Version 3.0/4.0
; SEQ ID NO 136
; LENGTH: 882
; TYPE: DNA
; ORGANISM: Chlamydia
; US-09-556-877-136

Query Match          22.2%; Score 24.6; DB 4; Length 882;
Best Local Similarity 76.9%; Pred. No. 12;
Matches 30; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

OY 8 GATCTTTTGCCCTTGACGAGATCTTTTCATCTTTCGACG 46
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 367 GATCTTTCACCTTGACGAGATCTTCTCTCTTTGCGGG 329

RESULT 14
US-09-620-412C-136/c
; Sequence 136, Application US/09620412C
; Patent No. 6448234
; GENERAL INFORMATION:
; APPLICANT: Steven P. Fling
; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATMENT AND
; FILE REFERENCE: 210121.469C7
; CURRENT APPLICATION NUMBER: US/09/620,412C
; CURRENT FILING DATE: 2000-07-20
; NUMBER OF SEQ ID NOS: 363
; SOFTWARE: FastSeq for Windows Version 3.0/4.0
; SEQ ID NO 136
; LENGTH: 882
; TYPE: DNA
; ORGANISM: Chlamydia
; US-09-620-412C-136

Query Match          22.2%; Score 24.6; DB 4; Length 882;
Best Local Similarity 76.9%; Pred. No. 12;
Matches 30; Conservative 0; Mismatches 9; Indels 0; Gaps 0;

OY 8 GATCTTTTGCCCTTGACGAGATCTTTTCATCTTTCGACG 46
   ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 367 GATCTTTCACCTTGACGAGATCTTCTCTCTTTGCGGG 329

RESULT 15
US-09-370-807-7
; Sequence 7, Application US/09370807
; Patent No. 6297034
; GENERAL INFORMATION:
; APPLICANT: Cahoon, Rebecca E.
; APPLICANT: Falco, S. Carl
; APPLICANT: Rafalski, J. Antoni
; APPLICANT: Sakai, Hajime
; TITLE OF INVENTION: N-End Rule Pathway Enzymes
; FILE REFERENCE: BB-1199
; CURRENT APPLICATION NUMBER: US/09/370,807
; CURRENT FILING DATE: 1999-08-09
; EARLIER APPLICATION NUMBER: 60/096,225
; EARLIER FILING DATE: August 12, 1998
; NUMBER OF SEQ ID NOS: 16
; SOFTWARE: Microsoft Office 97
; SEQ ID NO 7
; LENGTH: 2407
; TYPE: DNA
; ORGANISM: Oryza sativa
; US-09-370-807-7

Query Match          22.2%; Score 24.6; DB 4; Length 2407;
Best Local Similarity 57.0%; Pred. No. 17;
Matches 45; Conservative 0; Mismatches 34; Indels 0; Gaps 0;

OY 30 TTTTATCTTTTGACGAGACTTCTGGGCGCGAGTAGTAATAACTCTGGCTCTGTGT 89
   ||||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
```

Db 882 TTTTACAGCACTGCATTTGAATTCAGAACAACAGCAATTAGACTGCTGCTTTAAACA 941
Oy 90 GTGCTGAGTGGCTGCTCT 108
| | | | | | | | | |
Db 942 TTCTCTGATTAAGTGTGT 960

Search completed: April 25, 2003, 02:29:59
Job time : 67 secs

GenCore version 5.1.4.p5.4578
Copyright (c) 1993 - 2003 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2003, 02:28:45 ; Search time 69 Seconds
(without alignments)
1750.468 Million cell updates/sec

Title: US-09-513-999c-3792_COPY_51_161

Perfect score: 111
Sequence: 1 atgggttgatcttgcctt.....gctctgagtgctgctact 111

Scoring table: IDENTITY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 709820 seqs, 544064369 residues

1 number of hits satisfying chosen parameters: 1419640

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications_NA:*
1: /cgn2_6/ptodata/1/pubpna/US07_PUBCOMB.seq:*
2: /cgn2_6/ptodata/1/pubpna/PTCT_NEW_PUB.seq:*
3: /cgn2_6/ptodata/1/pubpna/US06_NEW_PUB.seq:*
4: /cgn2_6/ptodata/1/pubpna/US06_PUBCOMB.seq:*
5: /cgn2_6/ptodata/1/pubpna/US07_NEW_PUB.seq:*
6: /cgn2_6/ptodata/1/pubpna/PTCTUS_PUBCOMB.seq:*
7: /cgn2_6/ptodata/1/pubpna/US08_NEW_PUB.seq:*
8: /cgn2_6/ptodata/1/pubpna/US08_PUBCOMB.seq:*
9: /cgn2_6/ptodata/1/pubpna/US09_NEW_PUB.seq:*
10: /cgn2_6/ptodata/1/pubpna/US09_PUBCOMB.seq:*
11: /cgn2_6/ptodata/1/pubpna/US10_NEW_PUB.seq:*
12: /cgn2_6/ptodata/1/pubpna/US10_PUBCOMB.seq:*
13: /cgn2_6/ptodata/1/pubpna/US60_NEW_PUB.seq:*
14: /cgn2_6/ptodata/1/pubpna/US60_PUBCOMB.seq:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	58.2	52.4	570	US-09-864-761-9118	Sequence 9118, App
2	57.6	51.9	684973	US-09-263-959-1	Sequence 1, App11
3	52.8	47.6	428	US-09-918-993-8153	Sequence 8153, App
4	38.8	35.0	162	US-09-864-761-31393	Sequence 31393, A
5	38.8	35.0	519	US-09-864-761-14866	Sequence 14866, A
6	29.2	26.3	369	US-09-954-456-922	Sequence 922, App
7	29.2	26.3	1314	US-09-954-456-1538	Sequence 1538, App
8	29.2	26.3	1314	US-10-125-540-609	Sequence 609, App
9	29.2	26.3	1314	US-09-764-870-609	Sequence 609, App
10	28	25.2	342	US-10-125-540-118	Sequence 118, App
11	28	25.2	342	US-09-764-870-118	Sequence 118, App
12	27.2	24.5	412	US-09-783-590-1450	Sequence 1450, App
13	27.2	24.5	504	US-10-092-154-1665	Sequence 1665, App
14	27.2	24.5	504	US-10-092-154-1665	Sequence 1665, App
15	27.2	24.5	504	US-10-092-154-1665	Sequence 1665, App
16	27.2	24.5	504	US-09-764-847-1665	Sequence 1665, App
17	27.2	24.5	504	US-09-764-847-1665	Sequence 1665, App
18	27.2	24.5	504	US-09-764-847-1665	Sequence 1665, App
19	27	24.3	487	US-09-747-155-270	Sequence 270, App

20	26.6	24.0	6372	US-09-880-107-3948	Sequence 3948, App
21	26.4	23.8	385	US-09-770-791-90	Sequence 90, App1
22	26.4	23.8	1666	US-09-938-842A-966	Sequence 966, App
23	26.4	23.8	170834	US-09-835-232-7	Sequence 7, App11
24	26.2	23.6	26225	US-10-091-504-1276	Sequence 1276, App
25	26.2	23.6	26225	US-09-764-889-254	Sequence 254, App
26	26	23.4	430	US-09-876-889-254	Sequence 254, App
27	26	23.4	1548	US-09-925-302-211	Sequence 211, App
28	26	23.4	1620	US-09-954-456-325	Sequence 325, App
29	26	23.4	5641	US-10-015-219-1733	Sequence 1733, App
30	26	23.4	5709	US-10-015-219-1734	Sequence 1734, App
31	25.8	23.2	590	US-09-864-761-12014	Sequence 12014, A
32	25.8	23.2	1663	US-09-822-830A-348	Sequence 348, App
33	25.6	23.1	440	US-10-092-154-366	Sequence 366, App
34	25.6	23.1	440	US-09-764-847-366	Sequence 366, App
35	25.6	23.1	442	US-09-880-107-1917	Sequence 1917, App
36	25.6	23.1	1184	US-09-969-347-210	Sequence 210, App
37	25.6	23.1	32169	US-10-092-154-1963	Sequence 1963, App
38	25.6	23.1	32169	US-09-764-847-1963	Sequence 1963, App
39	25.4	22.9	11617	US-09-860-670-265	Sequence 265, App
40	25.2	22.7	390	US-09-770-791-11	Sequence 11, App1
41	25.2	22.7	455	US-09-960-352-9392	Sequence 9392, App
42	25.2	22.7	509	US-09-992-598-149	Sequence 149, App
43	25.2	22.7	509	US-09-989-735-149	Sequence 149, App
44	25.2	22.7	509	US-09-989-735-149	Sequence 149, App
45	25.2	22.7	509	US-09-990-444-149	Sequence 149, App

ALIGNMENTS

RESULT 1
US-09-864-761-9118/c
Sequence 9118, Application US/09864761
Patient No. US20020048763A1
GENERAL INFORMATION:
APPLICANT: Penn, Sharon G.
APPLICANT: Rank, David R.
APPLICANT: Hanzel, David K.
APPLICANT: Chen, Wensheng
TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
FILE REFERENCE: Aeomica-X-1
CURRENT APPLICATION NUMBER: US/09/864,761
CURRENT FILING DATE: 2001-05-23
PRIOR APPLICATION NUMBER: US 60/180,312
PRIOR FILING DATE: 2000-02-04
PRIOR APPLICATION NUMBER: US 60/207,456
PRIOR FILING DATE: 2000-05-26
PRIOR APPLICATION NUMBER: US 09/632,366
PRIOR FILING DATE: 2000-08-03
PRIOR APPLICATION NUMBER: GB 24263.6
PRIOR FILING DATE: 2000-10-04
PRIOR APPLICATION NUMBER: US 60/236,359
PRIOR FILING DATE: 2000-09-27
PRIOR APPLICATION NUMBER: PCT/US01/00666
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00667
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00664
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00669
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00665
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00668
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00663
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00662
PRIOR FILING DATE: 2001-01-30
PRIOR APPLICATION NUMBER: PCT/US01/00661
PRIOR FILING DATE: 2001-01-30

```

PRIORITY APPLICATION NUMBER: PCT/US01/00670
PRIORITY FILING DATE: 2001-01-30
PRIORITY APPLICATION NUMBER: US 60/234,687
PRIORITY FILING DATE: 2000-09-21
PRIORITY APPLICATION NUMBER: US 09/608,408
PRIORITY FILING DATE: 2000-06-30
PRIORITY APPLICATION NUMBER: US 09/774,203
PRIORITY FILING DATE: 2001-01-29
NUMBER OF SEQ. ID NOS: 49117
SOFTWARE: Annonmax Sequence Listing Engine vers. 1.1
SEQ. ID NO. 9118
LENGTH: 570
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
OTHER INFORMATION: MAP TO AP000053.1
OTHER INFORMATION: EXPRESSED IN HELIX, SIGNAL = 2.8
OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 2.4
OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 4.8
OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 4
OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 2.3
OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 3
OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 3.6
OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 4.4
US-09-864-761-9118

```

Query Match	52.48;	Score 58.2;	DB 10;	Length 570;
Best Local Similarity	79.38;	Pred. No. 1.2e-10;		
Matches 69; Conservative	0;	Mismatches 18;	Indels 0;	Gaps 0

Qy 24 GGATTCCTTTTTCATCCTTGACAGGACTCTGGGGCCGGAGATGTAAACCTCGGGTCT 83
| | | | | | | | | | | | | | | | | | | | | |
Db 566 GAATGATCTCCTGCCTTCGTGGGATTCTTGGCTGGAGATGTAAATTCCTGGGCT 507

```

Oy      84 CTGTGTGCTGCCTGAGTGGCTGCTTAC 110
          |||||
Db      506 TTGTGTGTGCCCTGAGTGGCGGCTCTGC 480

```

RESULT 2
US-09-263-959-1/c

Sequence 1, Application US/09263959
Patent No. US20020150891A1
GENERAL INFORMATION:
APPLICANT: HOOD, Leroy E.
APPLICANT: Rowen, Lee
APPLICANT: KOOP, Ben F.
TITLE OF INVENTION: DIAGNOSTIC AND THERAPEUTIC COMPOSITIONS AND METHODS WHICH UTILIZE
NUMBER OF SEQUENCES: 1279
CORRESPONDENCE ADDRESS:

ADDRESS: Seed and Berry LLP
STREET: 6300 Columbia Center, 701 Fifth Avenue
CITY: Seattle
STATE: Washington

```

; COUNTRY: US
; ZIP: 98104-7092
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy
; COVERED BY: TND PC

```

```

;
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
;
; 89-06-07 09:55:00

```

APPLICATION NUMBER: 05/0
FILING DATE: 05-MAR-1999
CLASSIFICATION:
ATTORNEY/AGENT INFORMATION

NAME: McMASTERS, DAVID D.
REGISTRATION NUMBER: 33,963
REFERENCE/DOCKET NUMBER: 920010.426C2
TELECOMMUNICATION INFORMATION:
TELEPHONE: (706) 622-4900

TELEPHONE: (206) 682-6031
TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 1:

```

; SEQUENCE CHARACTERISTICS:
;     LENGTH: 684973 base pairs
;     TYPE: nucleic acid
;     STRANDEDNESS: single
;     TOPOLOGY: linear
;
US-09-263-959-1

```

Query Match	51.9%;	Score 57.6;	DB 10;	Length 684973;
Best Local Similarity	78.4%;	Pred. No. 1.5e-09;		
Matches	69;	Conservative	0;	Mismatches 19;
			Indels	0;
			Gaps	0;

Qy 24 GGATCTTTTCATCTTTCGAGGGACTCTGGGCCGGAGTATGTAAACCTCCTGGTCT 83
||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 404732 GGACAGATTTCCTGCCTTGCTGGGGATCTGAGGCTGGAATGTGTAAAACTCCCTGGGTCT 404673

QY	84	CTGTGTGTCCTGAGTGCGCTCTACT	111
Db	404672	CTGTGTGTCCTGAGCAGCTGCTCTGCT	404645

RESULT 3
US-09-918-995-8153

? Sequence 8153, Application 05/09918995
 ? Publication No. US20030073623A1
 ? GENERAL INFORMATION:
 ? APPLICANT: Hyseng, Inc.
 ? TITLE OF INVENTION: NOVEL NUCLEIC ACID SEQUENCES OBTAINED
 ? TITLE OF INVENTION: FROM VARIOUS CDNA LIBRARIES
 ? FILE REFERENCE: 20411-756
 ? CURRENT APPLICATION NUMBER: US/09/918, 995
 ? CURRENT FILING DATE: 2001-07-30

```

; PRIOR APPLICATION NUMBER: US/09/235,076
; PRIOR FILING DATE: 1999-01-20
; NUMBER OF SEQ ID NOS: 38054
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 8153

```

```

; LENGTH: 428
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-918-995-8153

```

Query Match	47.6%;	Score 52.8;	DB 9;	Length 428;
Best Local Similarity	71.9%;	Pred. No. 8.2e-09;		
Matches	69;	Conservative	0;	Mismatches 27;
			Indels	0;
			Gaps	0

QY 15 TGCC^TTCGAGCATCTTTTCACTCTTGCGGGACTCTGGGGCCGAGTAGTAAACT 74
||||||| || ||| |||| | | | ||| ||||| ||| ||
Db 272 TGCCTTGCCTGAGATGCCATCACCCTTCTTTGGGGATCCTAAGGCTGGAGTAGTAAACGT 337

QY	75	CCTGGGTCCTGTGTGTGCTGAGTGCCTACTAC	110
Db	332	CCTGGGTCCTGTATGTGCTGAGCACCGTTC	367

RESULT 4
US-09-864-761-31393

```

: Sequence 31393, Application US/09864761
: Patent No. US20020048763A1
:
: GENERAL INFORMATION:
:
: APPLICANT: Pean, Sharron G.
: APPLICANT: Rank, David R.
: APPLICANT: Hanzel, David K.
: APPLICANT: Chen, Wensheng
:
: TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
: TITLE OF INVENTION: GENE EXPRESSION ANALYSIS BY MICROARRAY

```

```

; FILE REFERENCE: Aeomica-X-1
;
; CURRENT APPLICATION NUMBER: US/09/864,761
; CURRENT FILING DATE: 2001-05-23
;
; PRIOR APPLICATION NUMBER: US 60/180 312

```

7 PAVAR AFFILIATION NUMBER: US 00/180,511
 6 PRIOR FILING DATE: 2000-02-04
 5 PRIOR APPLICATION NUMBER: US 60/207,456
 4 PRIOR FILING DATE: 2000-05-26
 3 PRIOR APPLICATION NUMBER: US 09/632,366
 2
 1


```
;; PRIOR FILING DATE: 2000-08-03
;; PRIOR APPLICATION NUMBER: GB 24263.6
;; PRIOR FILING DATE: 2000-10-04
;; PRIOR APPLICATION NUMBER: US 60/236,359
;; PRIOR FILING DATE: 2000-09-27
;; PRIOR APPLICATION NUMBER: PCT/US01/00666
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00667
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00664
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00669
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00665
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00668
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00663
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00662
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00661
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00670
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: US 60/234,687
;; PRIOR FILING DATE: 2000-09-21
;; PRIOR APPLICATION NUMBER: US 09/608,408
;; PRIOR FILING DATE: 2000-06-30
;; PRIOR APPLICATION NUMBER: US 09/774,203
;; PRIOR FILING DATE: 2001-01-29
;; NUMBER OF SEQ ID NOS: 49117
;; SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
;; SEQ ID NO 31393
;; LENGTH: 162
;; TYPE: DNA
;; ORGANISM: Homo sapiens
;; FEATURE:
;; OTHER INFORMATION: MAP TO AC017089.2
;; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.8
;; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
;; OTHER INFORMATION: SWISSPROT HIT: P98161, EVALU6 3.00e+00
;; OTHER INFORMATION: EST_HUMAN HIT: AI79250.1, EVALU6 6.00e-07
;; OTHER INFORMATION: NT HIT: AL163210.2, EVALU6 4.00e-04
US-09-864-761-31393

Query Match          35.0%; Score 38.8; DB 10; Length 162;
Best Local Similarity 70.3%; Pred. No. 0.00043;
Matches 52; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

QY 38 CTTTGACAGGACTTCTGGGCGGAGATGTAAACTCTGTGCTGTGCTCTGA 97
    ||||| ||| ||| ||||| ||| ||||| ||| ||| ||||| |||||
Db 52 CTTTCCGTGAGTCCAGGCGCCAGAAATATCTAAGCTCTGAGATTTCATGTGCTGA 111
QY 98 GTGGCTGCTCTACT 111
    | | | | | | | | | |
Db 112 GCAGATGCTCTGCT 125

RESULT 5
US-09-864-761-14866
;; Sequence 14866, Application US/09864761
;; Patent No. US20020048763A1
;; GENERAL INFORMATION:
;; APPLICANT: Penn, Sharon G.
;; APPLICANT: Rank, David R.
;; APPLICANT: Hanzel, David K.
;; APPLICANT: Chen, Wensheng
;; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR
;; FILE REFERENCE: Aecomica-X-1
;; CURRENT APPLICATION NUMBER: US/09/864,761
;; CURRENT FILING DATE: 2001-05-23
```

```
;; PRIOR APPLICATION NUMBER: US 60/180,312
;; PRIOR FILING DATE: 2000-02-04
;; PRIOR APPLICATION NUMBER: US 60/207,456
;; PRIOR FILING DATE: 2000-05-26
;; PRIOR APPLICATION NUMBER: US 09/632,366
;; PRIOR FILING DATE: 2000-08-03
;; PRIOR APPLICATION NUMBER: GB 24263.6
;; PRIOR FILING DATE: 2000-10-04
;; PRIOR APPLICATION NUMBER: US 60/236,359
;; PRIOR FILING DATE: 2000-09-27
;; PRIOR APPLICATION NUMBER: PCT/US01/00666
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00667
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00664
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00669
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00665
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00668
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00663
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00662
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00661
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: PCT/US01/00670
;; PRIOR FILING DATE: 2001-01-30
;; PRIOR APPLICATION NUMBER: US 60/234,687
;; PRIOR FILING DATE: 2000-09-21
;; PRIOR APPLICATION NUMBER: US 09/608,408
;; PRIOR FILING DATE: 2000-06-30
;; PRIOR APPLICATION NUMBER: US 09/774,203
;; PRIOR FILING DATE: 2001-01-29
;; NUMBER OF SEQ ID NOS: 49117
;; SOFTWARE: Anomax Sequence Listing Engine vers. 1.1
;; SEQ ID NO 14866
;; LENGTH: 519
;; TYPE: DNA
;; ORGANISM: Homo sapiens
;; FEATURE:
;; OTHER INFORMATION: MAP TO AC017089.2
;; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.8
;; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 3.6
US-09-864-761-14866

Query Match          35.0%; Score 38.8; DB 10; Length 519;
Best Local Similarity 70.3%; Pred. No. 0.0006;
Matches 52; Conservative 0; Mismatches 22; Indels 0; Gaps 0;

QY 38 CTTTGACAGGACTTCTGGGCGGAGATGTAAACTCTGTGCTGTGCTCTGA 97
    ||||| ||| ||| ||||| ||| ||||| ||| ||| ||||| |||||
Db 301 CTTTCCGTGAGTCCAGGCGCCAGAAATATCTAAGCTCTGAGATTTCATGTGCTCTGA 360
QY 98 GTGGCTGCTCTACT 111
    | | | | | | | | | |
Db 361 GCAGATGCTCTGCT 374

RESULT 6
US-09-954-456-922/c
;; Sequence 922, Application US/09954456
;; Patent No. US20020115057A1
;; GENERAL INFORMATION:
;; APPLICANT: Young, Paul
;; TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents Using C
;; FILE REFERENCE: 689290-76
;; CURRENT APPLICATION NUMBER: US/09/954,456
;; CURRENT FILING DATE: 2001-09-18
;; PRIOR APPLICATION NUMBER: US/60/233,617
```

```

; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,052
; PRIOR FILING DATE: 2000-09-20
; PRIOR APPLICATION NUMBER: US/60/234,923
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,134
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,637
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,638
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,711
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,720
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,840
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,863
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 2276
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 922
; LENGTH: 369
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-954-456-922

Query Match          26.3%; Score 29.2; DB 10; Length 369;
Best Local Similarity 57.8%; Pred. No. 1.1;
Matches 52; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 1 ATGGGTGATCTTTGGCTTGCAGAGATCTTTTCATCTTTGCAGGACTTCTGGGCCG 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 143 AGGGCTGGAATTTAGTTTCAATATAGCTTCACGCTTAGCAATTAACCTAGTCCAA 84
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 61 GAGTATGTAACCTCTGGCTCTGTGTG 90
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 83 GACAATATTGATTCCTAGTTCTGTGG 54
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 7
US-09-954-456-1538/c
; Sequence 1538, Application US/09954456
; Patent No. US20020115057A1
; GENERAL INFORMATION:
; APPLICANT: Young, Paul
; TITLE OF INVENTION: Process for Identifying Anti-Cancer Therapeutic Agents using Candi-
; TITLE OF INVENTION: Sets
; FILE REFERENCE: 689290-76
; CURRENT APPLICATION NUMBER: US/09/954,456
; PRIOR FILING DATE: 2001-09-18
; PRIOR APPLICATION NUMBER: US/60/233,617
; PRIOR FILING DATE: 2000-09-18
; PRIOR APPLICATION NUMBER: US/60/234,052
; PRIOR FILING DATE: 2000-09-20
; PRIOR APPLICATION NUMBER: US/60/234,923
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,134
; PRIOR FILING DATE: 2000-09-25
; PRIOR APPLICATION NUMBER: US/60/235,637
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,638
; PRIOR FILING DATE: 2000-09-26
; PRIOR APPLICATION NUMBER: US/60/235,711
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,720
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,840
; PRIOR FILING DATE: 2000-09-27
; PRIOR APPLICATION NUMBER: US/60/235,863
; PRIOR FILING DATE: 2000-09-27
; NUMBER OF SEQ ID NOS: 2276
; SOFTWARE: PatentIn version 3.0
```

```

; SEQ ID NO 1538
; LENGTH: 369
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-954-456-1538

Query Match          26.3%; Score 29.2; DB 10; Length 369;
Best Local Similarity 57.8%; Pred. No. 1.1;
Matches 52; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 1 ATGGGTGATCTTTGGCTTGCAGAGATCTTTTCATCTTTGCAGGACTTCTGGGCCG 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 143 AGGGCTGGAATTTAGTTTCAATATAGCTTCACGCTTAGCAATTAACCTAGTCCAA 84
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 61 GAGTATGTAACCTCTGGCTCTGTGTG 90
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 83 GACAATATTGATTCCTAGTTCTGTGG 54
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 8
US-10-125-540-609
; Sequence 609, Application US/10125540
; Publication No. US20030059875A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT214C1
; CURRENT APPLICATION NUMBER: US/10/125,540
; PRIOR FILING DATE: 2002-04-19
; PRIOR APPLICATION REMOVED - See File Wrapper or Palm
; NUMBER OF SEQ ID NOS: 646
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 609
; LENGTH: 1314
; TYPE: DNA
; ORGANISM: Homo sapiens
US-10-125-540-609

Query Match          26.3%; Score 29.2; DB 9; Length 1314;
Best Local Similarity 56.1%; Pred. No. 1.6;
Matches 55; Conservative 0; Mismatches 43; Indels 0; Gaps 0;

QY 3 GGGTGAATCTTTGGCTTGCAGAGATCTTTTCATCTTTGCAGGACTTCTGGGCCG 62
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 982 GGGGTGAACAGTAGGCTGGGAGAAATTCCTCTGACAAAGAGTTGGAGATGAT 1041
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 63 GTATGTAACCTCTGGCTCTGTGTG 100
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 1042 GAATACAAACCCCTGGGACACTGCTGGCCCATAGTG 1079
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 9
US-09-764-870-609
; Sequence 609, Application US/09764870
; Patent No. US20020042386A1
; GENERAL INFORMATION:
; APPLICANT: Rosen et al.
; TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
; FILE REFERENCE: PT214
; CURRENT APPLICATION NUMBER: US/09/764,870
; PRIOR FILING DATE: 2001-01-17
; PRIOR APPLICATION DATA REMOVED - consult PALM or file wrapper
; NUMBER OF SEQ ID NOS: 646
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 609
; LENGTH: 1314
; TYPE: DNA
; ORGANISM: Homo sapiens
US-09-764-870-609

Query Match          26.3%; Score 29.2; DB 10; Length 1314;
Best Local Similarity 56.1%; Pred. No. 1.6;
Matches 55; Conservative 0; Mismatches 43; Indels 0; Gaps 0;
```

```
QY 3 GCGTGAATCTTTGCTTCGAGATCTTTTCATCTTTGACAGGACCTTCGTGGGCGGA 62
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 982 GGGGTGAACAGTAGCCCTGGGGGAGAAATGCTTCCTACAGAAAGATGGAGATGAT 1041
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 63 GTATGTAAACTCTGTGGTCTCTGTGTGTCCTGAGTG 100
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 1042 GAATACAAACCCTGGGACACTGCTGGCCCATAGTG 1079
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
```

RESULT 10

```
US-10-125-540-118
: Sequence 118, Application US/10125540
: Publication No. US20030059875A1
: GENERAL INFORMATION:
: APPLICANT: Rosen et al.
: TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
: FILE REFERENCE: PT214C1
: CURRENT APPLICATION NUMBER: US/10/125,540
: PRIOR FILING DATE: 2002-04-19
: Prior Application removed - See File Wrapper or Palm
: NUMBER OF SEQ ID NOS: 646
: SOFTWARE: Patentln Ver. 2.0
: SEQ ID NO 118
: LENGTH: 342
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (134)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (286)
: OTHER INFORMATION: n equals a,t,g, or c
US-10-125-540-118
```

```
Query Match 25.28; Score 28; DB 9; Length 342;
Best Local Similarity 55.08; Pred. No. 2.9;
Matches 55; Conservative 0; Mismatches 45; Indels 0; Gaps 0;
```

```
QY 1 ATGGGTGATCTTTGCTTCGAGATCTTTTCATCTTTGACAGGACCTTCGTGGGCGG 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 9 ACGAGGTGAACAGTAGCCCTGGGGGAGAAATGCTTCCTACAGAAAGATGGAGATG 68
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 61 GAGTATGTAACACTCTGTGGTCTCTGTGTGTCCTGAGTG 100
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
    69 ATGATACAAACCCTGGGACACTGCTGGCCCATAGTG 108
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
```

RESULT 11

```
US-09-764-870-118
: Sequence 118, Application US/09764870
: Patent No. US20020042386A1
: GENERAL INFORMATION:
: APPLICANT: Rosen et al.
: TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies
: FILE REFERENCE: PT214
: CURRENT APPLICATION NUMBER: US/09/764,870
: PRIOR FILING DATE: 2001-01-17
: Prior Application data removed - consult PALM or file wrapper
: NUMBER OF SEQ ID NOS: 646
: SOFTWARE: Patentln Ver. 2.0
: SEQ ID NO 118
: LENGTH: 342
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: SITE
: LOCATION: (134)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: SITE
: LOCATION: (286)
```

```
: OTHER INFORMATION: n equals a,t,g, or c
US-09-764-870-118
```

```
Query Match 25.28; Score 28; DB 10; Length 342;
Best Local Similarity 55.08; Pred. No. 2.9;
Matches 55; Conservative 0; Mismatches 45; Indels 0; Gaps 0;
```

```
QY 1 ATGGGTGATCTTTGCTTCGAGATCTTTTCATCTTTGACAGGACCTTCGTGGGCGG 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 9 ACGAGGTGAACAGTAGCCCTGGGGGAGAAATGCTTCCTACAGAAAGATGGAGATG 68
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
QY 61 GAGTATGTAACACTCTGTGGTCTCTGTGTGTCCTGAGTG 100
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 69 ATGATACAAACCCTGGGACACTGCTGGCCCATAGTG 108
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
```

RESULT 12

```
US-09-783-590-1450
: Sequence 1450, Application US/09783590
: Patent No. US20020110850A1
: GENERAL INFORMATION:
: APPLICANT: Dillon, Patrick J.
: APPLICANT: Haseltine, William A.
: APPLICANT: Li, Haodong
: APPLICANT: Rosen, Craig A.
: APPLICANT: Ruben, Steven M.
: TITLE OF INVENTION: Human Genes, Sequences, and Expression Products 16.2
: FILE REFERENCE: PO-16/2C1
: CURRENT APPLICATION NUMBER: US/09/783,590
: CURRENT FILING DATE: 2000-02-15
: PRIOR APPLICATION NUMBER: 08/420,856
: PRIOR FILING DATE: 1995-04-12
: PRIOR APPLICATION NUMBER: 08/346,731
: PRIOR FILING DATE: 1994-11-21
: NUMBER OF SEQ ID NOS: 12485
: SOFTWARE: Patentln Ver. 2.0
: SEQ ID NO 1450
: LENGTH: 412
: TYPE: DNA
: ORGANISM: Homo sapiens
: FEATURE:
: NAME/KEY: misc_feature
: LOCATION: (5)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (189)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (190)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (231)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (265)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (266)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (268)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (269)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (274)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (275)
: OTHER INFORMATION: n equals a,t,g, or c
: NAME/KEY: misc_feature
: LOCATION: (339)
```

OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (340)
OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (370)
OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (399)
OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (402)
OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (403)
OTHER INFORMATION: n equals a,t,g, or c
NAME/KEY: misc feature
LOCATION: (406)
OTHER INFORMATION: n equals a,t,g, or c
US-09-783-590-1450

Query Match 25.2%; Score 28; DB 10; Length 412;
Best Local Similarity 67.3%; Pred. No. 3;

Matches 37; Conservative 0; Mismatches 18; Indels 0; Gaps 0;

QY 57 GCCGAGATATGTAACCTCTGGGCTCTGTGTGCTGCTGAGTGGCTGCTCTACT 111
DB 213 GCCAGAGATCTGAGGCTTTGAACTGTGTGACAGGCCCTCAGTGGCTCTTNT 267

RESULT 13

US-10-092-154-1665/c

Sequence 1665, Application US/10092154

Publication No. US20030054375A1

GENERAL INFORMATION:

APPLICANT: Rosen et al.

TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies

FILE REFERENCE: PC009C1

CURRENT APPLICATION NUMBER: US/10/092,154

CURRENT FILING DATE: 2002-03-07

NUMBER OF SEQ ID NOS: 2003

Prior Application removed - See File Wrapper or Palm

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 1665

LENGTH: 504

TYPE: DNA

ORGANISM: Homo sapiens

US-10-092-154-1665

Query Match 24.5%; Score 27.2; DB 9; Length 504;
Best Local Similarity 56.8%; Pred. No. 6.1;

Matches 50; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 6 TGGATCTTTGCTTGCAGATTTCTTTTCATCTTTTCAGAGGACTTGGGCGGAGTA 65
DB 321 TTGGTTGTTTGTCTTCAGTCCTCTCTTTCTTAACACAGAGCCAGCTCTGCAATAGGC 262
QY 66 TGTAAACTCCTGGGCTCTGTGTGTGC 93
DB 261 AGGAACATGCTAGTCCAGTGTGTGC 234

RESULT 14

US-10-092-154-1666/c

Sequence 1666, Application US/10092154

Publication No. US20030054375A1

GENERAL INFORMATION:

APPLICANT: Rosen et al.

TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies

FILE REFERENCE: PC009C1

CURRENT APPLICATION NUMBER: US/10/092,154

CURRENT FILING DATE: 2002-03-07

NUMBER OF SEQ ID NOS: 2003

Prior Application removed - See File Wrapper or Palm
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 1666
LENGTH: 504
TYPE: DNA
ORGANISM: Homo sapiens
US-10-092-154-1666

Query Match 24.5%; Score 27.2; DB 9; Length 504;
Best Local Similarity 56.8%; Pred. No. 6.1;

Matches 50; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 6 TGGATCTTTGCTTGCAGATTTCTTTTCATCTTTTCAGAGGACTTGGGCGGAGTA 65
DB 321 TTGGTTGTTTGTCTTCAGTCCTCTCTTTCTTAACACAGAGCCAGCTCTGCAATAGGC 262
QY 66 TGTAAACTCCTGGGCTCTGTGTGTGC 93
DB 261 AGGAACATGCTAGTCCAGTGTGTGC 234

RESULT 15

US-10-092-154-1667/c

Sequence 1667, Application US/10092154

Publication No. US20030054375A1

GENERAL INFORMATION:

APPLICANT: Rosen et al.

TITLE OF INVENTION: Nucleic Acids, Proteins, and Antibodies

FILE REFERENCE: PC009C1

CURRENT APPLICATION NUMBER: US/10/092,154

CURRENT FILING DATE: 2002-03-07

NUMBER OF SEQ ID NOS: 2003

Prior Application removed - See File Wrapper or Palm

SOFTWARE: PatentIn Ver. 2.0

SEQ ID NO 1667

LENGTH: 504

TYPE: DNA

ORGANISM: Homo sapiens

US-10-092-154-1667

Query Match 24.5%; Score 27.2; DB 9; Length 504;
Best Local Similarity 56.8%; Pred. No. 6.1;

Matches 50; Conservative 0; Mismatches 38; Indels 0; Gaps 0;

QY 6 TGGATCTTTGCTTGCAGATTTCTTTTCATCTTTTCAGAGGACTTGGGCGGAGTA 65
DB 321 TTGGTTGTTTGTCTTCAGTCCTCTCTTTCTTAACACAGAGCCAGCTCTGCAATAGGC 262
QY 66 TGTAAACTCCTGGGCTCTGTGTGTGC 93
DB 261 AGGAACATGCTAGTCCAGTGTGTGC 234

Search completed: April 25, 2003, 03:07:48
Job time : 228 secs